

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: October 28, 2003, 15:13:35 ; Search time 1515 Seconds
(without alignments)
270.031 Million cell updates/sec

Title: US-09-335-032-71
Perfect score: 10
Sequence: 1 cttctctttt 10

Scoring table: OLIGO NUC
Gapop 60.0 , Gapext 60.0

Searched: 2888711 seqs, 2045481386 residues

Word size : 0

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 500 summaries

Database : GenEmbl.*

- 1: gb.ba.*
- 2: gb.htg.*
- 3: gb.in.*
- 4: gb.om.*
- 5: gb.ov.*
- 6: gb.pat.*
- 7: gb.ph.*
- 8: gb.pl.*
- 9: gb.pr.*
- 10: gb.ro.*
- 11: gb.sts.*
- 12: gb.sy.*
- 13: gb.un.*
- 14: gb.vi.*
- 15: em.ba.*
- 16: em.fun.*
- 17: em.hum.*
- 18: em.in.*
- 19: em.mu.*
- 20: em.om.*
- 21: em.or.*
- 22: em.ov.*
- 23: em.pat.*
- 24: em.ph.*
- 25: em.pl.*
- 26: em.ro.*
- 27: em.sts.*
- 28: em.un.*
- 29: em.vi.*
- 30: em.htg.hum.*
- 31: em.htg.inv.*
- 32: em.htg.other.*
- 33: em.htg.mus.*
- 34: em.htg.pln.*
- 35: em.htg.rod.*
- 36: em.htg.man.*
- 37: em.htg.vrt.*
- 38: em.sy.*
- 39: em.htgo.hum.*
- 40: em.htgo.mus.*
- 41: em.htgo.other.*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	10	100.0	10	6	BD065135	Character
C 2	10	100.0	12	6	AR029992	Sequence
C 3	10	100.0	12	6	AR030057	Sequence
C 4	10	100.0	17	6	AX500413	Sequence
C 5	10	100.0	17	6	AX500414	Sequence
C 6	10	100.0	17	6	AX500415	Sequence
C 7	10	100.0	17	6	AX500416	Sequence
C 8	10	100.0	17	6	AX500417	Sequence
C 9	10	100.0	17	6	AX500418	Sequence
C 10	10	100.0	17	6	AX500419	Sequence
C 11	10	100.0	17	6	AX500420	Sequence
C 12	10	100.0	17	6	AX673153	Sequence
C 13	10	100.0	19	6	AX659410	Sequence
C 14	10	100.0	20	6	AR315845	Sequence
C 15	10	100.0	20	6	I48976	Sequence
C 16	10	100.0	21	6	AR054595	Sequence
C 17	10	100.0	21	6	AR136775	Sequence
C 18	10	100.0	21	6	E35992	Sequence
C 19	10	100.0	22	6	AR029876	Sequence
C 20	10	100.0	22	6	AX352316	Sequence
C 21	10	100.0	22	6	AX352317	Sequence
C 22	10	100.0	22	6	AX352318	Sequence
C 23	10	100.0	24	6	AR069183	Sequence
C 24	10	100.0	24	6	AR102693	Sequence
C 25	10	100.0	24	6	AR175558	Sequence
C 26	10	100.0	24	6	AX443310	Sequence
C 27	10	100.0	24	6	BD090389	Sequence
C 28	10	100.0	24	6	BD176467	Sequence
C 29	10	100.0	24	6	I64401	Sequence
C 30	10	100.0	25	6	AX502410	Sequence
C 31	10	100.0	25	6	AX502411	Sequence
C 32	10	100.0	25	6	AX502412	Sequence
C 33	10	100.0	25	6	AX502413	Sequence
C 34	10	100.0	25	6	AX502414	Sequence
C 35	10	100.0	25	6	AX502415	Sequence
C 36	10	100.0	25	6	AX502416	Sequence
C 37	10	100.0	25	6	AX502417	Sequence
C 38	10	100.0	25	6	AX502418	Sequence
C 39	10	100.0	25	6	AX502419	Sequence
C 40	10	100.0	25	6	AX502420	Sequence
C 41	10	100.0	25	6	AX502421	Sequence
C 42	10	100.0	25	6	AX502422	Sequence
C 43	10	100.0	25	6	AX502423	Sequence
C 44	10	100.0	25	6	AX502424	Sequence
C 45	10	100.0	25	6	AX502425	Sequence
C 46	10	100.0	25	6	I48977	Sequence
C 47	10	100.0	26	8	ATH525492	Arabidops
C 48	10	100.0	26	8	ATH525503	Arabidops
C 49	10	100.0	27	6	AR066258	Sequence
C 50	10	100.0	27	6	AR191561	Sequence
C 51	10	100.0	27	6	AX115839	Sequence
C 52	10	100.0	28	6	AX040823	Sequence
C 53	10	100.0	29	6	AR261655	Sequence
C 54	10	100.0	30	6	AR261654	Sequence
C 55	10	100.0	30	6	AR261656	Sequence
C 56	10	100.0	30	6	AR261657	Sequence
C 57	10	100.0	33	6	AR120483	Sequence
C 58	10	100.0	33	6	BD063492	Sequence
C 59	10	100.0	33	6	E36416	Sequence
C 60	10	100.0	38	6	AX218433	Sequence
C 61	10	100.0	38	6	AX220343	Sequence
C 62	10	100.0	38	6	AX273375	Sequence
C 63	10	100.0	40	6	BD182426	Sequence
C 64	10	100.0	42	6	I12092	Sequence
C 65	10	100.0	43	6	AX484623	Sequence

C 66	10	100.0	45	6	AX061875	AX061875 Sequence	C 139	10	100.0	111	11	BX295946	BX295946 Arabidops
C 67	10	100.0	47	8	ATH528602	AJ258602 Arabidops	C 140	10	100.0	112	9	HS1087R	Z56291 H. sapiens C
C 68	10	100.0	47	6	AR284713	AR284713 Sequence	C 141	10	100.0	113	8	ATH526836	AJ526836 Arabidops
C 69	10	100.0	47	6	AR288787	AR288787 Sequence	C 142	10	100.0	113	8	ATH526837	AJ526837 Arabidops
C 70	10	100.0	47	6	AR289547	AR289547 Sequence	C 143	10	100.0	116	9	HS197G4R	Z55121 H. sapiens C
C 71	10	100.0	47	6	AR289811	AR289811 Sequence	C 144	10	100.0	117	9	HS68D7R	Z65938 H. sapiens C
C 72	10	100.0	47	6	AR290737	AR290737 Sequence	C 145	10	100.0	117	9	HS68E6F	Z55835 H. sapiens C
C 73	10	100.0	47	6	AR291549	AR291549 Sequence	C 146	10	100.0	118	6	AX554775	AX554775 Sequence
C 74	10	100.0	47	6	AR139658	AR139658 Sequence	C 147	10	100.0	118	11	HSPF39G05	AL009929 H. sapiens
C 75	10	100.0	50	6	AX158156	AX158156 Sequence	C 148	10	100.0	120	11	HUMIFARE	ML1287 Human alpha
C 76	10	100.0	51	6	AX118141	AX118141 Sequence	C 149	10	100.0	120	11	HUMSWX752	L24631 Human chrom
C 77	10	100.0	51	6	AX158155	AX158155 Sequence	C 150	10	100.0	121	6	AX262806	AX262806 Sequence
C 78	10	100.0	51	6	AX160381	AX160381 Sequence	C 151	10	100.0	121	6	AX262807	AX262807 Sequence
C 79	10	100.0	51	6	AX160382	AX160382 Sequence	C 152	10	100.0	121	6	AX262822	AX262822 Sequence
C 80	10	100.0	51	6	AX160383	AX160383 Sequence	C 153	10	100.0	121	6	AX262823	AX262823 Sequence
C 81	10	100.0	51	6	AX165562	AX165562 Sequence	C 154	10	100.0	121	6	AX262926	AX262926 Sequence
C 82	10	100.0	51	8	ATH521150	AJ521150 Arabidops	C 155	10	100.0	121	6	AX262927	AX262927 Sequence
C 83	10	100.0	53	6	AF098682	AF098682 Sequence	C 156	10	100.0	121	6	AX262930	AX262930 Sequence
C 84	10	100.0	53	6	AR098683	AR098683 Sequence	C 157	10	100.0	121	6	AX262931	AX262931 Sequence
C 85	10	100.0	53	6	AR204756	AR204756 Sequence	C 158	10	100.0	121	8	AY201631	AY201631 Arabidops
C 86	10	100.0	53	6	AR204757	AR204757 Sequence	C 159	10	100.0	121	11	G32506	G32506 A003F19 Hum
C 87	10	100.0	54	6	AR134108	AR134108 Sequence	C 160	10	100.0	122	11	G43213	G43213 WIAF-1715-S
C 88	10	100.0	54	6	AR134285	AR134285 Sequence	C 161	10	100.0	122	11	G43214	G43214 WIAF-1716-S
C 89	10	100.0	57	8	ATH505724	AJ505724 Arabidops	C 162	10	100.0	125	6	BD033545	BD033545 Sequence
C 90	10	100.0	58	8	ATH527067	AJ527067 Arabidops	C 163	10	100.0	125	6	BD033545	BD033545 Sequence
C 91	10	100.0	61	6	AX270701	AX270701 Sequence	C 164	10	100.0	128	1	HEA71U	M33452 H. influenza
C 92	10	100.0	61	6	AX272232	AX272232 Sequence	C 165	10	100.0	128	4	OAU35059	U35059 Ovis aries
C 93	10	100.0	65	6	AX482835	AX482835 Sequence	C 166	10	100.0	129	11	G32708	G32708 A009027 Hum
C 94	10	100.0	65	6	AX482852	AX482852 Sequence	C 167	10	100.0	130	8	ATH505656	AJ505656 Arabidops
C 95	10	100.0	65	6	AX485490	AX485490 Sequence	C 168	10	100.0	130	8	ATH528046	AJ528046 Arabidops
C 96	10	100.0	70	6	AX1656	AX1656 Sequence 1	C 169	10	100.0	130	9	HS191F9F	Z57722 H. sapiens C
C 97	10	100.0	70	6	AX207788	AX207788 Sequence	C 170	10	100.0	130	11	G18970	Z65038 H. sapiens C
C 98	10	100.0	71	6	I62432	I62432 Sequence 5	C 171	10	100.0	130	11	G18970	G18970 cow STS BMS
C 99	10	100.0	71	9	AH006998S06	AF026858 Homo sapi	C 172	10	100.0	131	11	HUMSWX53	L14998 Human chrom
C 100	10	100.0	73	6	BD055583	BD055583 Sequence	C 173	10	100.0	134	9	HSPAI3A8	Z78634 H. sapiens f
C 101	10	100.0	75	6	I62433	I62433 Sequence 6	C 174	10	100.0	135	8	AF503161	AF503161 Ceiba pen
C 102	10	100.0	76	6	AX463283	AX463283 Sequence	C 175	10	100.0	135	10	MUSNR2C06	L35019 Mouse N-net
C 103	10	100.0	76	6	BD038896	BD038896 Sequence	C 176	10	100.0	136	6	AX135409	AX135409 Sequence
C 104	10	100.0	78	6	I62446	I62446 Sequence 19	C 177	10	100.0	136	11	HSPF56A10	AL034100 H. sapiens
C 105	10	100.0	80	11	BX000934	BX000934 Arabidops	C 178	10	100.0	138	6	BD048886	BD048886 Sequence
C 106	10	100.0	81	14	AB015322	AB015322 Hepatitis	C 179	10	100.0	138	6	BD061135	BD061135 Secreted
C 107	10	100.0	81	14	AF148872	AF148872 Norwalk-1	C 180	10	100.0	138	9	HSPAI5A11	Z78713 H. sapiens f
C 108	10	100.0	81	14	AF148873	AF148873 Norwalk-1	C 181	10	100.0	138	11	HUMUT799A	L39147 Human STS U
C 109	10	100.0	82	6	I62447	I62447 Sequence 20	C 182	10	100.0	140	9	HUMHELAD	D45429 Homo sapien
C 110	10	100.0	83	6	AX386001	AX386001 Sequence	C 183	10	100.0	140	11	G20401	G20401 human STS A
C 111	10	100.0	83	6	BD037643	BD037643 Sequence	C 184	10	100.0	141	6	BD055897	BD055897 Sequence
C 112	10	100.0	83	6	BD112411	BD112411 EST and e	C 185	10	100.0	141	11	AU046839	AU046839 Rattus no
C 113	10	100.0	86	6	AX240905	AX240905 Sequence	C 186	10	100.0	145	9	AY190094	AY190094 Homo sapi
C 114	10	100.0	86	11	DM173H11S	Z32462 D. melanoga	C 187	10	100.0	146	6	AR139657	AR139657 Sequence
C 115	10	100.0	88	11	BX295958	BX295958 Arabidops	C 188	10	100.0	146	11	G02302	G02302 human STS S
C 116	10	100.0	88	11	G66316	G66316 sy2268 YAC	C 189	10	100.0	147	3	AF320603	AF320603 Trigon a c
C 117	10	100.0	91	3	DME426842	AJ426842 Drosophil	C 190	10	100.0	147	6	AX379108	AX379108 Sequence
C 118	10	100.0	93	6	AR126785	AR126785 Sequence	C 191	10	100.0	147	6	BD046874	BD046874 Sequence
C 119	10	100.0	93	6	AR202442	AR202442 Sequence	C 192	10	100.0	147	9	AF515842	AF515842 Homo sapi
C 120	10	100.0	93	6	AX522948	AX522948 Sequence	C 193	10	100.0	148	6	BD043676	BD043676 Sequence
C 121	10	100.0	95	6	BD118772	BD118772 EST and e	C 194	10	100.0	148	6	BD043896	BD043896 Sequence
C 122	10	100.0	98	8	YSCPIPR4	ML1711 yeast RNA p	C 195	10	100.0	148	9	AY190090	AY190090 Homo sapi
C 123	10	100.0	98	10	RNPCTP5	AF040266 Rattus no	C 196	10	100.0	148	9	HUMFCGAA02	M90722 Human FC-ga
C 124	10	100.0	99	6	BD038421	BD038421 Sequence	C 197	10	100.0	149	6	AX259898	AX259898 Sequence
C 125	10	100.0	99	6	BD038552	BD038552 Sequence	C 198	10	100.0	149	9	HS9F11R	Z61827 H. sapiens C
C 126	10	100.0	102	11	G20394	G20394 human STS A	C 199	10	100.0	150	11	G59542	G59542 SHGC-130099
C 127	10	100.0	103	8	OSA532497	AJ532497 Oryza sat	C 200	10	100.0	151	9	G59725	G59725 SHGC-130478
C 128	10	100.0	103	9	HSSTEE4TF1	X84366 H. sapiens t	C 201	10	100.0	151	11	AY190102	AY190102 Homo sapi
C 129	10	100.0	103	11	G19611	G19611 human STS A	C 202	10	100.0	152	6	BD028580	BD028580 Sequence
C 130	10	100.0	104	6	AX645544	AX645544 Sequence	C 203	10	100.0	152	9	AY190114	AY190114 Homo sapi
C 131	10	100.0	104	6	AX676705	AX676705 Sequence	C 204	10	100.0	152	9	HUMALURPTF	M37552 Human AFP g
C 132	10	100.0	104	6	BD049479	BD049479 Sequence	C 205	10	100.0	153	5	AY269268	AY269268 Gila eleg
C 133	10	100.0	104	8	AY220694	AY220694 Oryza sat	C 206	10	100.0	153	5	AX269268	AX269268 H. sapiens m
C 134	10	100.0	105	6	AX127420	AX127420 Sequence	C 207	10	100.0	153	11	G64362	G64362 P316119/T7
C 135	10	100.0	105	11	BX248221	BX248221 Arabidops	C 208	10	100.0	156	6	AX435756	AX435756 Sequence
C 136	10	100.0	105	11	G71016	G71016 A09121234FB	C 209	10	100.0	156	11	G30715	AJ276202 Homo sapi
C 137	10	100.0	107	9	HUMHELAC	D45428 Homo sapien	C 210	10	100.0	156	11	HSTEB0301	X85348 H. sapiens t
C 138	10	100.0	110	11	BX295578	BX295578 Arabidops	C 211	10	100.0	157	6	AX196537	AX196537 Sequence

212	10	100.0	10	100.0	10	100.0	285	194	11	GI2787	GI2787 sWS2144 Er
213	10	100.0	10	100.0	10	100.0	286	195	6	BD050159	Sequence
c 214	10	100.0	10	100.0	10	100.0	c 287	195	11	EX293660	Arabidops
215	10	100.0	10	100.0	10	100.0	288	195	11	EX293660	Arabidops
216	10	100.0	10	100.0	10	100.0	289	196	11	ECU35065	U35065 Escherichia
217	10	100.0	10	100.0	10	100.0	c 290	196	9	H87B3F	Z63570 H.sapiens C
218	10	100.0	10	100.0	10	100.0	291	196	9	H87B3R	Z63571 H.sapiens C
219	10	100.0	10	100.0	10	100.0	c 292	196	11	EX294583	Arabidops
220	10	100.0	10	100.0	10	100.0	293	197	6	AX186932	Sequence
c 221	10	100.0	10	100.0	10	100.0	c 294	197	6	AX186932	Sequence
c 222	10	100.0	10	100.0	10	100.0	c 295	198	6	AX202139	Sequence
c 223	10	100.0	10	100.0	10	100.0	c 296	198	6	AX311050	Sequence
224	10	100.0	10	100.0	10	100.0	c 297	198	9	HSA405781	BD030797 Homo sapi
c 225	10	100.0	10	100.0	10	100.0	c 298	199	6	BD030797	Sequence
c 226	10	100.0	10	100.0	10	100.0	c 299	199	9	HS167F1R	Z53637 H.sapiens C
227	10	100.0	10	100.0	10	100.0	300	200	11	BV012598	Sequence
c 228	10	100.0	10	100.0	10	100.0	301	201	6	AX173101	BV012598 M6S151 Hu
c 229	10	100.0	10	100.0	10	100.0	c 302	201	6	AX276551	Sequence
230	10	100.0	10	100.0	10	100.0	c 303	201	6	AX435807	Sequence
231	10	100.0	10	100.0	10	100.0	304	201	6	BD040764	Sequence
232	10	100.0	10	100.0	10	100.0	c 305	201	9	HSA405663	Homo sapi
233	10	100.0	10	100.0	10	100.0	c 306	201	9	HSA405788	Homo sapi
c 234	10	100.0	10	100.0	10	100.0	c 307	202	5	AY029202	AY029202 Gallus ga
c 235	10	100.0	10	100.0	10	100.0	c 308	203	3	AY185121	AY185121 Leishmani
c 236	10	100.0	10	100.0	10	100.0	309	203	3	LSP270148	AY270148 Leishmani
c 237	10	100.0	10	100.0	10	100.0	c 310	203	9	AF262665	AF262665 Homo sapi
c 238	10	100.0	10	100.0	10	100.0	c 311	203	11	G20532	G20532 human STS A
c 239	10	100.0	10	100.0	10	100.0	c 312	204	6	BD042756	BD042756 Sequence
c 240	10	100.0	10	100.0	10	100.0	c 313	204	9	HSA405760	AY405760 Homo sapi
c 241	10	100.0	10	100.0	10	100.0	c 314	204	9	HSA405762	AY405762 Homo sapi
c 242	10	100.0	10	100.0	10	100.0	c 315	204	10	WMIG7P1A	X55507 M.musculus
c 243	10	100.0	10	100.0	10	100.0	c 316	204	11	G20705	X55507 human STS A
c 244	10	100.0	10	100.0	10	100.0	c 317	205	6	AX387628	G20705 human STS A
c 245	10	100.0	10	100.0	10	100.0	c 318	205	9	HUNGMP1405	AX387628 Sequence
c 246	10	100.0	10	100.0	10	100.0	319	206	6	AR251645	M60223 Human granu
c 247	10	100.0	10	100.0	10	100.0	320	206	8	ATH551904	AR251645 Sequence
c 248	10	100.0	10	100.0	10	100.0	c 321	206	10	AF361008	AF361008 Mus muscu
249	10	100.0	10	100.0	10	100.0	322	207	8	AB043755	AB043755 Fagus cre
c 250	10	100.0	10	100.0	10	100.0	323	207	8	AB043756	AB043756 Fagus cre
c 251	10	100.0	10	100.0	10	100.0	c 324	207	9	HSAA011594	AB043756 Fagus cre
c 252	10	100.0	10	100.0	10	100.0	c 325	207	9	HSAA011594	AB043756 Fagus cre
c 253	10	100.0	10	100.0	10	100.0	326	208	6	AX310754	AB043756 Fagus cre
c 254	10	100.0	10	100.0	10	100.0	327	208	6	AF242801	AX310754 Homo sapi
c 255	10	100.0	10	100.0	10	100.0	c 328	208	11	AF235213	AX310754 Homo sapi
c 256	10	100.0	10	100.0	10	100.0	329	209	3	LSP270149	AF235213 Sus scrof
c 257	10	100.0	10	100.0	10	100.0	c 330	210	6	AX415513	AX270149 Leishmani
c 258	10	100.0	10	100.0	10	100.0	c 331	210	6	AX431292	AX415513 Sequence
c 259	10	100.0	10	100.0	10	100.0	c 332	210	6	AX431292	AX431292 Sequence
c 260	10	100.0	10	100.0	10	100.0	333	210	6	AX482093	AX431292 Sequence
c 261	10	100.0	10	100.0	10	100.0	334	212	6	AX511332	AX482093 Sequence
c 262	10	100.0	10	100.0	10	100.0	335	212	6	AX721693	AX511332 Sequence
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c 264	10	100.0	10	100.0	10	100.0	337	212	6	BD045031	BD045031 Sequence
c 265	10	100.0	10	100.0	10	100.0	338	212	6	BD028873	U27369 Human Menke
c 266	10	100.0	10	100.0	10	100.0	c 339	213	6	BD053618	BD028873 Sequence
c 267	10	100.0	10	100.0	10	100.0	c 340	213	9	HSAA05752	BD053618 Sequence
c 268	10	100.0	10	100.0	10	100.0	c 341	213	9	HSAA05752	BD053618 Sequence
c 269	10	100.0	10	100.0	10	100.0	c 342	214	5	WSP428343	AX405752 Homo sapi
c 270	10	100.0	10	100.0	10	100.0	343	215	1	RP282717	AX405752 Homo sapi
c 271	10	100.0	10	100.0	10	100.0	344	215	1	WSP428343	AX405752 Homo sapi
c 272	10	100.0	10	100.0	10	100.0	c 345	216	3	CECBH11H	AF114764 Ictallurus
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ALIGNMENTS


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LOCUS      BD065135
DEFINITION Characterization of the yeast transcriptome.
ACCESSION  BD065135
VERSION     BD065135.1 GI:22610738
KEYWORDS   JP 2001509017-A/71.
SOURCE     Saccharomyces cerevisiae (baker's yeast)
ORGANISM   Saccharomycetes cerevisiae
REFERENCE  1 (bases 1 to 10)
AUTHORS    Wang, C.-G. and Hepburn, A.G.
TITLE      Genetic sequence assay using DNA triple strand formation
JOURNAL    Patent: US 5861244-A 71 10-JUL-2001;
           THE JOHNS HOPKINS UNIVERSITY SCHOOL OF MEDICINE
COMMENT    OS Saccharomyces cerevisiae (yeast)
           PN JP 2001509017-A/71
           PD 10-JUL-2001
           PF 22-JAN-1998 JP 1998532117
           PR 23-JAN-1997 US 60/035917
           PI VICTOR E VELCULESCU, BERT VOGELSTEIN, KENNETH W KINZLER PC
           CI 2N15/10, CI 2N15/31, C07K14/395, CI 2Q1/68, CI 2Q1/02 CC
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DEFINITION Sequence 181 from patent US 5861244.
ACCESSION  AR029992
VERSION     AR029992.1 GI:5943206
KEYWORDS   Unknown.
SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 12)
AUTHORS    Wang, C.-G. and Hepburn, A.G.
TITLE      Genetic sequence assay using DNA triple strand formation
JOURNAL    Patent: US 5861244-A 191 19-JAN-1999;
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LOCUS      BD065135
DEFINITION Characterization of the yeast transcriptome.
ACCESSION  BD065135
VERSION     BD065135.1 GI:22610738
KEYWORDS   JP 2001509017-A/71.
SOURCE     Saccharomyces cerevisiae (baker's yeast)
ORGANISM   Saccharomycetes cerevisiae
REFERENCE  1 (bases 1 to 12)
AUTHORS    Wang, C.-G. and Hepburn, A.G.
TITLE      Genetic sequence assay using DNA triple strand formation
JOURNAL    Patent: US 5861244-A 71 10-JUL-2001;
           THE JOHNS HOPKINS UNIVERSITY SCHOOL OF MEDICINE
COMMENT    OS Saccharomyces cerevisiae (yeast)
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           PI VICTOR E VELCULESCU, BERT VOGELSTEIN, KENNETH W KINZLER PC
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RESULT 4
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DEFINITION Sequence 1720 from Patent EP1229046.
ACCESSION  AX500413
VERSION     AX500413.1 GI:23382706
KEYWORDS   Homo sapiens (human)
SOURCE     Homo sapiens
ORGANISM   Homo sapiens
REFERENCE  1
AUTHORS    Zhan, J.
TITLE      Human testis expressed patched like protein
JOURNAL    Patent: EP 1229046-A 1720 07-AUG-2002;
           Aeomica, Inc. (US)
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RESULT 5
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DEFINITION Sequence 1721 from Patent EP1229046.
ACCESSION  AX500414
VERSION     AX500414.1 GI:23382707
KEYWORDS   Homo sapiens (human)
SOURCE     Homo sapiens
ORGANISM   Homo sapiens

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RESULT 3
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LOCUS      AR030057
DEFINITION Sequence 246 from patent US 5861244.
ACCESSION  AR030057
VERSION     AR030057.1 GI:5943271
KEYWORDS   Unknown.
SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 12)
AUTHORS    Wang, C.-G. and Hepburn, A.G.
TITLE      Genetic sequence assay using DNA triple strand formation
JOURNAL    Patent: US 5861244-A 246 19-JAN-1999;
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RESULT 4
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DEFINITION Sequence 1720 from Patent EP1229046.
ACCESSION  AX500413
VERSION     AX500413.1 GI:23382706
KEYWORDS   Homo sapiens (human)
SOURCE     Homo sapiens
ORGANISM   Homo sapiens
REFERENCE  1
AUTHORS    Zhan, J.
TITLE      Human testis expressed patched like protein
JOURNAL    Patent: EP 1229046-A 1720 07-AUG-2002;
           Aeomica, Inc. (US)
FEATURES   Location/Qualifiers
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RESULT 5
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LOCUS      AX500414
DEFINITION Sequence 1721 from Patent EP1229046.
ACCESSION  AX500414
VERSION     AX500414.1 GI:23382707
KEYWORDS   Homo sapiens (human)
SOURCE     Homo sapiens
ORGANISM   Homo sapiens

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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1
REFERENCE
AUTHORS      Zhan,J.
TITLE        Human testis expressed patched like protein
JOURNAL      Patent: EP 1229046-A 1721 07-AUG-2002;
              Aeomica, Inc. (US)
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DEFINITION   Sequence 1722 from Patent EP1229046.
ACCESSION    AX500415
VERSION      AX500415.1 GI:23382708
KEYWORDS     Homo sapiens (human)
ORGANISM     Homo sapiens
REFERENCE    1
AUTHORS      Zhan,J.
TITLE        Human testis expressed patched like protein
JOURNAL      Patent: EP 1229046-A 1722 07-AUG-2002;
              Aeomica, Inc. (US)
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DEFINITION   Sequence 1723 from Patent EP1229046.
ACCESSION    AX500416
VERSION      AX500416.1 GI:23382709
KEYWORDS     Homo sapiens (human)
ORGANISM     Homo sapiens
REFERENCE    1
AUTHORS      Zhan,J.
TITLE        Human testis expressed patched like protein
JOURNAL      Patent: EP 1229046-A 1723 07-AUG-2002;
              Aeomica, Inc. (US)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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AUTHORS      Zhan,J.
TITLE        Human testis expressed patched like protein
JOURNAL      Patent: EP 1229046-A 1721 07-AUG-2002;
              Aeomica, Inc. (US)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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AUTHORS      Zhan,J.
TITLE        Human testis expressed patched like protein
JOURNAL      Patent: EP 1229046-A 1721 07-AUG-2002;
              Aeomica, Inc. (US)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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AUTHORS      Zhan,J.
TITLE        Human testis expressed patched like protein
JOURNAL      Patent: EP 1229046-A 1724 07-AUG-2002;
              Aeomica, Inc. (US)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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REFERENCE
AUTHORS      Zhan,J.
TITLE        Human testis expressed patched like protein
JOURNAL      Patent: EP 1229046-A 1724 07-AUG-2002;
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DEFINITION   Sequence 1724 from Patent EP1229046.
ACCESSION    AX500417
VERSION      AX500417.1 GI:23382710
KEYWORDS     Homo sapiens (human)
ORGANISM     Homo sapiens
REFERENCE    1
AUTHORS      Zhan,J.
TITLE        Human testis expressed patched like protein
JOURNAL      Patent: EP 1229046-A 1724 07-AUG-2002;
              Aeomica, Inc. (US)
FEATURES
source       Location/Qualifiers
              1..17
              /organism="Homo sapiens"
              /mol_type="genomic DNA"
              /db_xref="taxon:9606"
BASE COUNT   8 a _ 0 c 6 g 3 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

RESULT 9
AX500418/c
LOCUS        AX500418      17 bp      DNA      linear      PAT 27-SEP-2002
DEFINITION   Sequence 1725 from Patent EP1229046.
ACCESSION    AX500418
VERSION      AX500418.1 GI:23382711
KEYWORDS     Homo sapiens (human)
ORGANISM     Homo sapiens
REFERENCE    1
AUTHORS      Zhan,J.
TITLE        Human testis expressed patched like protein
JOURNAL      Patent: EP 1229046-A 1725 07-AUG-2002;
              Aeomica, Inc. (US)
FEATURES
source       Location/Qualifiers
              1..17
              /organism="Homo sapiens"
              /mol_type="genomic DNA"
              /db_xref="taxon:9606"
BASE COUNT   8 a _ 1 c 5 g 3 t
ORIGIN

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Query Match      100.0%; Score 10; DB 6; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 12 CTTCTCTTTT 3

RESULT 10
AX500419/c
LOCUS      AX500419      17 bp      DNA      linear      PAT 27-SEP-2002
DEFINITION Sequence 1726 from Patent EP1229046.
ACCESSION  AX500419
VERSION     AX500419.1 GI:23382712
KEYWORDS
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
REFERENCE   1
AUTHORS     Zhan,J.
TITLE       Human testis expressed patched like protein
JOURNAL     Patent: EP 1229046-A 1726 07-AUG-2002;
            Acomica, Inc. (US)
FEATURES    source
            1..17
            /organism="Homo sapiens"
            /mol_type="genomic DNA"
            /db_xref="taxon:9606"
BASE COUNT  8 a 1 c 4 g 4 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 11 CTTCTCTTTT 2

RESULT 11
AX500420/c
LOCUS      AX500420      17 bp      DNA      linear      PAT 27-SEP-2002
DEFINITION Sequence 1727 from Patent EP1229046.
ACCESSION  AX500420
VERSION     AX500420.1 GI:23382713
KEYWORDS
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
REFERENCE   1
AUTHORS     Zhan,J.
TITLE       Human testis expressed patched like protein
JOURNAL     Patent: EP 1229046-A 1727 07-AUG-2002;
            Acomica, Inc. (US)
FEATURES    source
            1..17
            /organism="Homo sapiens"
            /mol_type="genomic DNA"
            /db_xref="taxon:9606"
BASE COUNT  8 a 1 c 5 g 3 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 1 CTTCTCTTTT 13

RESULT 14

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Db 10 CTTCTCTTTT 1

RESULT 12
AX673153
LOCUS      AX673153      17 bp      DNA      linear      PAT 27-MAR-2003
DEFINITION Sequence 1598 from Patent WO03004526.
ACCESSION  AX673153
VERSION     AX673153.1 GI:29331501
KEYWORDS
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
REFERENCE   1
AUTHORS     Telerman,A., Anson,R. and Tuijnder,M.
TITLE       Sequences involved in phenomena of tumour suppression, tumour
            reversion, apoptosis and/or resistance to viruses and their use as
            medicines
JOURNAL     Patent: WO 03004526-A 1598 16-JAN-2003;
            Molecular Engines Laboratories (FR)
FEATURES    source
            1..17
            /organism="Homo sapiens"
            /mol_type="genomic DNA"
            /db_xref="taxon:9606"
BASE COUNT  2 a 6 c 1 g 8 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 6 CTTCTCTTTT 15

RESULT 13
AX659410
LOCUS      AX659410      19 bp      DNA      linear      PAT 22-MAR-2003
DEFINITION Sequence 12 from Patent WO02102824.
ACCESSION  AX659410
VERSION     AX659410.1 GI:29161640
KEYWORDS
SOURCE      synthetic construct
            synthetic construct
            artificial sequences.
REFERENCE   1
AUTHORS     Beifohr,C. and Shaidr,J.
TITLE       Method for specific fast detection of relevant bacteria in drinking
            water
JOURNAL     Patent: WO 02102824-A 12 27-DEC-2002;
            Vermicon AG (DE)
FEATURES    Location/Qualifiers
            source
            1..19
            /organism="synthetic construct"
            /mol_type="genomic DNA"
            /db_xref="taxon:32630"
            /note="oligonucleotide"
BASE COUNT  1 a 7 c 2 g 9 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 19;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 4 CTTCTCTTTT 13

RESULT 14

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AR315845/c
LOCUS AR315845 20 bp DNA linear PAT 12-JUN-2003
DEFINITION Sequence 6382 from patent US 6559294.
ACCESSION AR315845
VERSION AR315845.1 GI:31709271
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 20)
AUTHORS Griffais, R., Hoigeth, S.K., Zagursky, R.J., Metcalf, B.J., Peek, J.A., Sankaran, B., and Fletcher, L.D.
TITLE Chlamydia pneumoniae polynucleotides and uses thereof
JOURNAL Patent: US 6559294-A 6382 06-MAY-2003;
FEATURES
Location/Qualifiers
1..20
/organism="unknown"
BASE COUNT 8 a 4 c 4 g 4 t
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 17 CTTCTCTTTT 8

RESULT 15
LOCUS I48976 20 bp DNA linear PAT 07-OCT-1997
DEFINITION Sequence 3 from patent US 5627054.
ACCESSION I48976
VERSION I48976.1 GI:2467439
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 20)
AUTHORS Gillespie, D. deceased.
TITLE Competitor primer asymmetric polymerase chain reaction
JOURNAL Patent: US 5627054-A 3 06-MAY-1997;
FEATURES
Location/Qualifiers
1..20
/organism="unknown"
BASE COUNT 4 a 5 c 3 g 8 t
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17

RESULT 16
LOCUS AR054595/c 21 bp DNA linear PAT 29-SEP-1999
DEFINITION Sequence 16 from patent US 5837447.
ACCESSION AR054595
VERSION AR054595.1 GI:5980172
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 21)
AUTHORS Gorski, J.
TITLE Monitoring an immune response by analysis of amplified immunoglobulin or T-cell-receptor nucleic acid
JOURNAL Patent: US 5837447-A 16 17-NOV-1998;

FEATURES
source Location/Qualifiers
1..21
/organism="unknown"
BASE COUNT 8 a 3 c 7 g 3 t
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

RESULT 17
LOCUS ARI36775/c 21 bp DNA linear PAT 16-JUN-2001
DEFINITION Sequence 8 from patent US 6162435.
ACCESSION ARI36775
VERSION ARI36775.1 GI:14478025
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 21)
AUTHORS Minion, F. Chris. and Hsu, T.
TITLE Recombinant mycoplasma hyopneumoniae vaccine
JOURNAL Patent: US 6162435-A 8 19-DEC-2000;
FEATURES
Location/Qualifiers
1..21
/organism="unknown"
BASE COUNT 12 a 0 c 7 g 2 t
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 15 CTTCTCTTTT 6

RESULT 18
LOCUS E35992/c 21 bp DNA linear PAT 31-JAN-2002
DEFINITION Method for detecting Kawasaki disease factor.
ACCESSION E35992
VERSION E35992.1 GI:18624703
KEYWORDS JP 2000157297-A/83.
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1 (bases 1 to 21)
AUTHORS Yoshioka, T. and Suzuki, R.
TITLE Method for detecting Kawasaki disease factor
JOURNAL Patent: JP 2000157297-A 83 13-JUN-2000;
COMMENT SHIONOGI & CO LTD
OS Artificial Sequence
PN JP 2000157297-A/83
PD 13-JUN-2000
PF 01-DEC-1998 JP 1998341661
PR TAKESHI YOSHIOKA, RYUJI SUZUKI
PI C12Q1/68, C12N15/09, G01N33/48, C12N15/00
CC CC
FH Key Location/Qualifiers
FT source 1..21
/organism='Artificial Sequence'.
FEATURES
source Location/Qualifiers
1..21
/organism="synthetic construct"

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/mol_type="genomic DNA"
/db_xref="taxon:32630"
3 t

BASE COUNT      8 a      3 c      7 g      3 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 18 CTTCTCTTTT 9

RESULT 19
AR029876
LOCUS      AR029876      22 bp      DNA      linear      PAT 29-SEP-1999
DEFINITION Sequence 65 from patent US 5861244.
ACCESSION AR029876
VERSION    AR029876.1 GI:5943090
KEYWORDS
SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 22)
AUTHORS    Wang, C.-G. and Hepburn, A.G.
TITLE      Genetic sequence assay using DNA triple strand formation
JOURNAL    Patent: US 5861244-A 65 19-JAN-1999;
FEATURES   Location/Qualifiers
            source
              1..22
              /organism="unknown"
BASE COUNT 1 a      5 c      0 g      16 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 22;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 6 CTTCTCTTTT 15

RESULT 20
AR029876
LOCUS      AR029876      22 bp      DNA      linear      PAT 06-FEB-2002
DEFINITION Sequence 612 from Patent WO0193902.
ACCESSION AR029876
VERSION    AR029876.1 GI:18617599
KEYWORDS
SOURCE     synthetic construct
ORGANISM   synthetic construct
REFERENCE  1
AUTHORS    Mond, J.J., Flora, M. and Klinman, D.M.
TITLE      Immunostimulatory rna/dna hybrid molecules
JOURNAL    Patent: WO 0193902-A 612 13-DEC-2001;
FEATURES   Location/Qualifiers
            source
              1..22
              /organism="synthetic construct"
              /mol_type="genomic DNA"
              /db_xref="taxon:32630"
              /note="Synthetic HDR"
BASE COUNT 1 a      4 c      3 g      14 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 22;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 1 CTTCTCTTTT 10

RESULT 21
AR029876
LOCUS      AR029876      22 bp      DNA      linear      PAT 06-FEB-2002
DEFINITION Sequence 613 from Patent WO0193902.
ACCESSION AR029876
VERSION    AR029876.1 GI:18617600
KEYWORDS
SOURCE     synthetic construct
ORGANISM   synthetic construct
REFERENCE  1
AUTHORS    Mond, J.J., Flora, M. and Klinman, D.M.
TITLE      Immunostimulatory rna/dna hybrid molecules
JOURNAL    Patent: WO 0193902-A 613 13-DEC-2001;
FEATURES   Location/Qualifiers
            source
              1..22
              /organism="synthetic construct"
              /mol_type="genomic DNA"
              /db_xref="taxon:32630"
              /note="Synthetic HDR"
BASE COUNT 1 a      4 c      4 g      13 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 22;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 7 CTTCTCTTTT 16

RESULT 22
AR029876
LOCUS      AR029876      22 bp      DNA      linear      PAT 06-FEB-2002
DEFINITION Sequence 614 from Patent WO0193902.
ACCESSION AR029876
VERSION    AR029876.1 GI:18617601
KEYWORDS
SOURCE     synthetic construct
ORGANISM   synthetic construct
REFERENCE  1
AUTHORS    Mond, J.J., Flora, M. and Klinman, D.M.
TITLE      Immunostimulatory rna/dna hybrid molecules
JOURNAL    Patent: WO 0193902-A 614 13-DEC-2001;
FEATURES   Location/Qualifiers
            source
              1..22
              /organism="synthetic construct"
              /mol_type="genomic DNA"
              /db_xref="taxon:32630"
              /note="Synthetic HDR"
BASE COUNT 1 a      4 c      2 g      15 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 22;
Best Local Similarity 100.0%; Pred. No. 1.4e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 7 CTTCTCTTTT 16

RESULT 23
AR069183/c
LOCUS      AR069183      24 bp      DNA      linear      PAT 18-FEB-2000
DEFINITION Sequence 17 from patent US 5891623.

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ACCESSION      AR069183
VERSION        AR069183.1  GI:7220071
KEYWORDS
SOURCE        Unknown.
ORGANISM      Unclassified.
REFERENCE     1 (bases 1 to 24)
AUTHORS      Primi,D.
TITLE        Diagnosis and treatment of AIDS onset
JOURNAL      Patent: US 5891623-A 17 06-APR-1999;
FEATURES     Location/Qualifiers
              1..24
              /organism="unknown"
BASE COUNT   10 a 3 c 7 g 4 t
ORIGIN
Query Match  100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
    |||||
Db 18 CTTCTCTTTT 9

RESULT 24
ACCESSION      AR102693
LOCUS          AR102693/c
DEFINITION     Sequence 16 from patent US 6087096.
ACCESSION      AR102693
VERSION        AR102693.1  GI:12814281
KEYWORDS
SOURCE        Unknown.
ORGANISM      Unclassified.
REFERENCE     1 (bases 1 to 24)
AUTHORS      Dau,P.C. and Liu,D.
TITLE        Method of intrafamily fragment analysis of the T cell receptor
              .alpha. and .beta. chain CDR3 regions
JOURNAL      Patent: US 6087096-A 16 11-JUL-2000;
FEATURES     Location/Qualifiers
              1..24
              /organism="unknown"
BASE COUNT   10 a 3 c 7 g 4 t
ORIGIN
Query Match  100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
    |||||
Db 18 CTTCTCTTTT 9

RESULT 25
ACCESSION      AR175558
LOCUS          AR175558/c
DEFINITION     Sequence 3 from patent US 6309837.
ACCESSION      AR175558
VERSION        AR175558.1  GI:117916857
KEYWORDS
SOURCE        Unknown.
ORGANISM      Unclassified.
REFERENCE     1 (bases 1 to 24)
AUTHORS      Dean,R.A. and Wang,Y.-H.
TITLE        PCR-based method for identifying a fusarium wilt-resistant genotype
              in plants
JOURNAL      Patent: US 6309837-A 3 30-OCT-2001;
FEATURES     Location/Qualifiers
              1..24
              /organism="unknown"
source

BASE COUNT   10 a 3 c 7 g 4 t
ORIGIN
Query Match  100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
    |||||
Db 18 CTTCTCTTTT 9

RESULT 26
ACCESSION      AX443310
LOCUS          AX443310/c
DEFINITION     Sequence 46 from Patent WO0216940.
ACCESSION      AX443310
VERSION        AX443310.1  GI:21690705
KEYWORDS
SOURCE        synthetic construct
              synthetic construct
              artificial sequences.
REFERENCE     1
AUTHORS      Sulavik,M., Ling,L.L., Opperman,T., Moir,D.T. and Bunker,C.
TITLE        Genomics-assisted rapid identification of targets
JOURNAL      Patent: WO 0216940-A 46 28-FEB-2002;
              Genome Therapeutics Corporation (US)
FEATURES     Location/Qualifiers
              1..24
              /organism="synthetic construct"
              /mol_type="genomic DNA"
              /db_xref="taxon:32630"
              /note="primer"
BASE COUNT   9 a 4 c 6 g 5 t
ORIGIN
Query Match  100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
    |||||
Db 16 CTTCTCTTTT 7

RESULT 27
ACCESSION      BD090389
LOCUS          BD090389/c
DEFINITION     A method of arraying genome clone.
ACCESSION      BD090389
VERSION        BD090389.1  GI:22635999
KEYWORDS
SOURCE        synthetic construct
              synthetic construct
              artificial sequences.
REFERENCE     1 (bases 1 to 24)
AUTHORS      Soeda,E.
TITLE        A method of arraying genome clone
JOURNAL      Patent: JP 2001321190-A 2633 20-NOV-2001;
              THE INSTITUTE OF PHYSICAL AND CHEMICAL RESEARCH, YUGENKAISHA
              GENOTECHS
COMMENT      OS Artificial Sequence
              PN JP 2001321190-A/2633
              PD 20-NOV-2001
              PF 12-MAR-2001 JP 2001068285
              PI EIICHI SOEDA
              PC C12N15/09,C12N15/09,C12M1/00,C12Q1/68,G01N33/53,G01N33/566, PC
              C12N15/00
              CC Description of Artificial Sequence:Synthetic DNA FH key
              Location/Qualifiers
              1..24
              /organism='Artificial Sequence'.
              FT source
              FT

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FEATURES
  source
    Location/Qualifiers
      1..24
      /organism="synthetic construct"
      /mol_type="genomic DNA"
      /db_xref="taxon:32630"
BASE COUNT      12 a      2 c      6 g      4 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY      1 CTCTCTCTTT 10
      |||
Db      19 CTCTCTCTTT 10
      |||

RESULT 28
BD176467/c
LOCUS      BD176467 24 bp DNA linear PAT 18-MAR-2003
DEFINITION      A method of arraying genome clone.
ACCESSION      BD176467
VERSION      BD176467.1 GI:29122175
KEYWORDS      WO 02072815-A/267.
SOURCE      synthetic construct
ORGANISM      artificial sequences.
REFERENCE      1 (bases 1 to 24)
AUTHORS      Soeda,E.
TITLE      A method of arraying genome clone
JOURNAL      Patent: WO 02072815-A 267 19-SEP-2002;
      EIICHI SOEDA,TAKESHI KUKITA
COMMENT      OS Artificial Sequence
      PN WO 02072815-A/267
      PD 19-SEP-2002 WO 2001JP004139
      PF 17-MAY-2001 WO 2001JP004139
      PR 12-MAR-2001 JP 01P 68285
      PT EIICHI SOEDA
      PC C12N15/09,C12Q1/68
      CC Description of Artificial Sequence: Synthetic DNA FH Key
      FT source
      FT      1..24
      Location/Qualifiers
        1..24
        /organism="synthetic construct"
        /mol_type="genomic DNA"
        /db_xref="taxon:32630"
BASE COUNT      12 a      2 c      6 g      4 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY      1 CTCTCTCTTT 10
      |||
Db      19 CTCTCTCTTT 10
      |||

RESULT 29
I64401/c
LOCUS      I64401 24 bp DNA linear PAT 07-OCT-1997
DEFINITION      Sequence 17 from patent US 5665355.
ACCESSION      I64401
VERSION      I64401.1 GI:2481295
KEYWORDS      .
SOURCE      Unknown.
ORGANISM      Unknown.
REFERENCE      1 (bases 1 to 24)
AUTHORS      Primi,D.
TITLE      Diagnosis and treatment of AIDS onset

JOURNAL Patent: US 5665355-A 17 09-SEP-1997;
  source
    Location/Qualifiers
      1..24
      /organism="unknown"
BASE COUNT      10 a      3 c      7 g      4 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY      1 CTCTCTCTTT 10
      |||
Db      18 CTCTCTCTTT 9
      |||

RESULT 30
AX502410/c
LOCUS      AX502410 25 bp DNA linear PAT 27-SEP-2002
DEFINITION      Sequence 3717 from Patent EP1229046.
ACCESSION      AX502410
VERSION      AX502410.1 GI:23384703
KEYWORDS      Homo sapiens (human)
SOURCE      Homo sapiens
ORGANISM      Homo sapiens
REFERENCE      1
AUTHORS      Zhan,J.
TITLE      Human testis expressed patched like protein
JOURNAL      Patent: EP 1229046-A 3717 07-AUG-2002;
      Aeomica, Inc. (US)
FEATURES
  source
    Location/Qualifiers
      1..25
      /organism="Homo sapiens"
      /mol_type="genomic DNA"
      /db_xref="taxon:9606"
BASE COUNT      9 a      2 c      9 g      5 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY      1 CTCTCTCTTT 10
      |||
Db      25 CTCTCTCTTT 16
      |||

RESULT 31
AX502411/c
LOCUS      AX502411 25 bp DNA linear PAT 27-SEP-2002
DEFINITION      Sequence 3718 from Patent EP1229046.
ACCESSION      AX502411
VERSION      AX502411.1 GI:23384704
KEYWORDS      Homo sapiens (human)
SOURCE      Homo sapiens
ORGANISM      Homo sapiens
REFERENCE      1
AUTHORS      Zhan,J.
TITLE      Human testis expressed patched like protein
JOURNAL      Patent: EP 1229046-A 3718 07-AUG-2002;
      Aeomica, Inc. (US)
FEATURES
  source
    Location/Qualifiers
      1..25
      /organism="Homo sapiens"
      /mol_type="genomic DNA"
      /db_xref="taxon:9606"
BASE COUNT      8 a      2 c      9 g      6 t
ORIGIN
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Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 24 CTTCTCTTTT 15

RESULT 32
AX502412/c
LOCUS
DEFINITION Sequence 3719 from Patent EP1229046.
ACCESSION AX502412
VERSION AX502412.1 GI:23384705
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1
AUTHORS Zhan, J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3719 07-AUG-2002;
Aeomica, Inc. (US)

FEATURES
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/mol_type="genomic DNA"
/db_xref="taxon:9606"

BASE COUNT 9 a 2 c 9 g 5 t
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Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
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Db 23 CTTCTCTTTT 14

RESULT 33
AX502413/c
LOCUS
DEFINITION Sequence 3720 from Patent EP1229046.
ACCESSION AX502413
VERSION AX502413.1 GI:23384706
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1
AUTHORS Zhan, J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3720 07-AUG-2002;
Aeomica, Inc. (US)

FEATURES
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BASE COUNT 9 a 2 c 9 g 5 t
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Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
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Qy 1 CTTCTCTTTT 10
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Db 22 CTTCTCTTTT 13

RESULT 34
AX502414/c
LOCUS
DEFINITION Sequence 3721 from Patent EP1229046.
ACCESSION AX502414
VERSION AX502414.1 GI:23384707
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1
AUTHORS Zhan, J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3721 07-AUG-2002;
Aeomica, Inc. (US)

FEATURES
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/mol_type="genomic DNA"
/db_xref="taxon:9606"

BASE COUNT 9 a 2 c 8 g 6 t
ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 21 CTTCTCTTTT 12

RESULT 35
AX502415/c
LOCUS
DEFINITION Sequence 3722 from Patent EP1229046.
ACCESSION AX502415
VERSION AX502415.1 GI:23384708
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1
AUTHORS Zhan, J.
TITLE Human testis expressed patched like protein
JOURNAL Patent: EP 1229046-A 3722 07-AUG-2002;
Aeomica, Inc. (US)

FEATURES
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Location/Qualifiers
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/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"

BASE COUNT 9 a 2 c 8 g 6 t
ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 20 CTTCTCTTTT 11

RESULT 36
AX502416/c
LOCUS
DEFINITION Sequence 3723 from Patent EP1229046.
ACCESSION AX502416

VERSION	AX502416.1	GI:23384709
SOURCE	Homo sapiens (human)	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	
AUTHORS	Zhan,J.	
TITLE	Human testis expressed patched like protein	
JOURNAL	Patent: EP 1229046-A 3723 07-AUG-2002;	
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BASE COUNT	9 a 1 c 8 g 7 t	
ORIGIN	1..25 /organism="Homo sapiens" /mol_type="genomic DNA" /db_xref="taxon:9606"	
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Matches	10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
Qy	1 CTTCTCTTTT 10	
Db	19 CTTCTCTTTT 10	
RESULT 37		
AX502417/c		PAT 27-SEP-2002
LOCUS	AX502417	25 bp DNA linear
DEFINITION	Sequence 3724 from Patent EP1229046.	
ACCESSION	AX502417	
VERSION	AX502417.1	GI:23384710
KEYWORDS	Homo sapiens (human)	
SOURCE	Homo sapiens	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	
AUTHORS	Zhan,J.	
TITLE	Human testis expressed patched like protein	
JOURNAL	Patent: EP 1229046-A 3724 07-AUG-2002;	
FEATURES	source	
BASE COUNT	9 a 1 c 9 g 6 t	
ORIGIN	1..25 /organism="Homo sapiens" /mol_type="genomic DNA" /db_xref="taxon:9606"	
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Best Local Similarity	100.0%; Pred. No. 1.3e+05;	
Matches	10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
Qy	1 CTTCTCTTTT 10	
Db	18 CTTCTCTTTT 9	
RESULT 38		
AX502418/c		PAT 27-SEP-2002
LOCUS	AX502418	25 bp DNA linear
DEFINITION	Sequence 3725 from Patent EP1229046.	
ACCESSION	AX502418	
VERSION	AX502418.1	GI:23384711
KEYWORDS	Homo sapiens (human)	
SOURCE	Homo sapiens	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	

AUTHORS	Zhan,J.	
TITLE	Human testis expressed patched like protein	
JOURNAL	Patent: EP 1229046-A 3725 07-AUG-2002;	
FEATURES	source	
BASE COUNT	9 a 2 c 9 g 5 t	
ORIGIN	1..25 /organism="Homo sapiens" /mol_type="genomic DNA" /db_xref="taxon:9606"	
Query Match	100.0%; Score 10; DB 6; Length 25;	
Best Local Similarity	100.0%; Pred. No. 1.3e+05;	
Matches	10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
Qy	1 CTTCTCTTTT 10	
Db	17 CTTCTCTTTT 8	
RESULT 39		
AX502419/c		PAT 27-SEP-2002
LOCUS	AX502419	25 bp DNA linear
DEFINITION	Sequence 3726 from Patent EP1229046.	
ACCESSION	AX502419	
VERSION	AX502419.1	GI:23384712
KEYWORDS	Homo sapiens (human)	
SOURCE	Homo sapiens	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	
AUTHORS	Zhan,J.	
TITLE	Human testis expressed patched like protein	
JOURNAL	Patent: EP 1229046-A 3726 07-AUG-2002;	
FEATURES	source	
BASE COUNT	9 a 3 c 8 g 5 t	
ORIGIN	1..25 /organism="Homo sapiens" /mol_type="genomic DNA" /db_xref="taxon:9606"	
Query Match	100.0%; Score 10; DB 6; Length 25;	
Best Local Similarity	100.0%; Pred. No. 1.3e+05;	
Matches	10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
Qy	1 CTTCTCTTTT 10	
Db	16 CTTCTCTTTT 7	
RESULT 40		
AX502420/c		PAT 27-SEP-2002
LOCUS	AX502420	25 bp DNA linear
DEFINITION	Sequence 3727 from Patent EP1229046.	
ACCESSION	AX502420	
VERSION	AX502420.1	GI:23384713
KEYWORDS	Homo sapiens (human)	
SOURCE	Homo sapiens	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	
AUTHORS	Zhan,J.	
TITLE	Human testis expressed patched like protein	
JOURNAL	Patent: EP 1229046-A 3727 07-AUG-2002;	
FEATURES	source	
BASE COUNT	9 a 3 c 8 g 5 t	
ORIGIN	1..25 /organism="Homo sapiens"	

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6 t
BASE COUNT
ORIGIN
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Query Match      100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 15 CTTCTCTTTT 6

RESULT 41
AX502421/c
LOCUS      AX502421      25 bp      DNA      linear      PAT 27-SEP-2002
DEFINITION Sequence 3728 from Patent EP1229046.
ACCESSION  AX502421
VERSION     AX502421.1 GI:23384714
KEYWORDS   .
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1
AUTHORS   Zhan,J.
TITLE     Human testis expressed patched like protein
JOURNAL   Patent: EP 1229046-A 3728 07-AUG-2002;
          Aeomica, Inc. (US)
FEATURES   Location/Qualifiers
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              /organism="Homo sapiens"
              /mol_type="genomic DNA"
              /db_xref="taxon:9606"
BASE COUNT      9 a      3 c      7 g
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 14 CTTCTCTTTT 5

RESULT 42
AX502422/c
LOCUS      AX502422      25 bp      DNA      linear      PAT 27-SEP-2002
DEFINITION Sequence 3729 from Patent EP1229046.
ACCESSION  AX502422
VERSION     AX502422.1 GI:23384715
KEYWORDS   .
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1
AUTHORS   Zhan,J.
TITLE     Human testis expressed patched like protein
JOURNAL   Patent: EP 1229046-A 3729 07-AUG-2002;
          Aeomica, Inc. (US)
FEATURES   Location/Qualifiers
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              /db_xref="taxon:9606"
BASE COUNT      9 a      3 c      7 g
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 11 CTTCTCTTTT 2

RESULT 43
AX502423/c
LOCUS      AX502423      25 bp      DNA      linear      PAT 27-SEP-2002
DEFINITION Sequence 3730 from Patent EP1229046.
ACCESSION  AX502423
VERSION     AX502423.1 GI:23384716
KEYWORDS   .
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1
AUTHORS   Zhan,J.
TITLE     Human testis expressed patched like protein
JOURNAL   Patent: EP 1229046-A 3730 07-AUG-2002;
          Aeomica, Inc. (US)
FEATURES   Location/Qualifiers
            source
              1..25
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              /mol_type="genomic DNA"
              /db_xref="taxon:9606"
BASE COUNT      9 a      3 c      7 g
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
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Db 12 CTTCTCTTTT 3

RESULT 44
AX502424/c
LOCUS      AX502424      25 bp      DNA      linear      PAT 27-SEP-2002
DEFINITION Sequence 3731 from Patent EP1229046.
ACCESSION  AX502424
VERSION     AX502424.1 GI:23384717
KEYWORDS   .
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1
AUTHORS   Zhan,J.
TITLE     Human testis expressed patched like protein
JOURNAL   Patent: EP 1229046-A 3731 07-AUG-2002;
          Aeomica, Inc. (US)
FEATURES   Location/Qualifiers
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              /db_xref="taxon:9606"
BASE COUNT      9 a      4 c      6 g
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
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Db 11 CTTCTCTTTT 2
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RESULT 45
AX502425/c
LOCUS AX502425 25 bp DNA linear PAT 27-SEP-2002
DEFINITION Sequence 3732 from Patent EP1229046.
ACCESSION AX502425
VERSION AX502425.1 GI:23384718
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 Zhan, J.
AUTHORS Human testis expressed patched like protein
TITLE Patent: EP 1229046-A 3732 07-AUG-2002;
JOURNAL Aeomica, Inc. (US)
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/organism="Homo sapiens"
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/db_xref="taxon:9606"
BASE COUNT 9 a 4 c 6 g 6 t
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1 CTTCTCTTTT 10
2 |||||
10 CTTCTCTTTT 1
148977
LOCUS I48977 25 bp DNA linear PAT 07-OCT-1997
DEFINITION Sequence 4 from patent US 5627054.
ACCESSION I48977
VERSION I48977.1 GI:2467440
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE
1 (bases 1 to 25)
AUTHORS Gillespie, D. deceased.
TITLE Competitor primer asymmetric polymerase chain reaction
JOURNAL Patent: US 5627054-A 4 06-MAY-1997;
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/organism="unknown"
BASE COUNT 4 a 5 c 3 g 8 t 5 others
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Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17

RESULT 47
ATH525492/c
LOCUS ATH525492 26 bp DNA linear PLN 29-MAR-2003
DEFINITION Arabidopsis thaliana T-DNA flanking sequence, left border, clone 097H03.
ACCESSION AJ525492
VERSION AJ525492.1 GI:26793728
KEYWORDS left border; T-DNA flanking sequence.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

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Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 Brunaud, V., Balzerque, S., Dubreucq, B., Aubourg, S., Samson, F.,
Chauvin, S., Bechtold, N., Cruaud, C., DeRose, R., Pelletier, G.,
Lepiniec, L., Caboche, M. and Lecharny, A.
T-DNA integration into the Arabidopsis genome depends on sequences
of pre-insertion sites
EMBO Rep. 3 (12), 1152-1157 (2002)
22363535
12446565
REFERENCE 2 (bases 1 to 26)
AUTHORS Balzerque, S.
TITLE Direct Submision
JOURNAL Submitted (21-NOV-2002) Balzerque S., UMRGV, INRA/CNRS, 2 rue
Gaston Cremieux, 91057 Evry cedex, FRANCE
COMMENT PCR was performed on DNA from transformants of Arabidopsis thaliana
plants from INRA (Versailles). The DNA fragment(s) resulting from
the PCR were directly sequenced from the left or the right border
to determine the genomic sequence flanking the insertion. T-DNA
derived sequences were removed. Information to order the
corresponding mutant line and a link to a database providing a
graphical display of the insertion site are available at
http://dbgap.versailles.inra.fr/publiclines/. This sequence has
been generated in the framework of the French plant genomics
program 'Genoplante' (http://www.genoplante.com and
http://genoplante-info.infobiogen.fr).
FEATURES
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/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/cultivar="Wassiliewskija"
/db_xref="taxon:3702"
/clone="097H03"
/clone_lib="Arabidopsis thaliana T-DNA insertion lines"
misc_feature
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/Note="T-DNA flanking sequence"
BASE COUNT 13 a 3 c 4 g 6 t
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Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 21 CTTCTCTTTT 12

RESULT 48
ATH525503/c
LOCUS ATH525503 26 bp DNA linear PLN 29-MAR-2003
DEFINITION Arabidopsis thaliana T-DNA flanking sequence, left border, clone 098B06.
ACCESSION AJ525503
VERSION AJ525503.1 GI:26793739
KEYWORDS left border; T-DNA flanking sequence.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 Brunaud, V., Balzerque, S., Dubreucq, B., Aubourg, S., Samson, F.,
Chauvin, S., Bechtold, N., Cruaud, C., DeRose, R., Pelletier, G.,
Lepiniec, L., Caboche, M. and Lecharny, A.
T-DNA integration into the Arabidopsis genome depends on sequences
of pre-insertion sites
EMBO Rep. 3 (12), 1152-1157 (2002)
22363535
12446565
REFERENCE 2 (bases 1 to 26)
AUTHORS
TITLE
JOURNAL
MEDLINE
PUBMED

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AUTHORS Balzergue,S.
 TITLE Direct Submission
 JOURNAL Submitted (21-NOV-2002) Balzergue S., UMRGV, INRA/CNRS, 2 rue Gaston Cremieux, 91057 Evry cedex, FRANCE

COMMENT PCR was performed on DNA from transformants of Arabidopsis thaliana plants from INRA (Versailles). The DNA fragment(s) resulting from the PCR were directly sequenced from the left or the right border to determine the genomic sequence flanking the insertion. T-DNA derived sequences were removed. Information to order the corresponding mutant line and a link to a database providing a graphical display of the insertion site are available at <http://dbsgap.versailles.inra.fr/publiclines/>. This sequence has been generated in the framework of the French plant genomics program "Genoplante" (<http://www.genoplante.com> and <http://genoplante-info.infobiogen.fr>).

FEATURES
 source
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 /organism="Arabidopsis thaliana"
 /mol_type="genomic DNA"
 /cultivar="Wassillewskija"
 /db_xref="taxon:3702"
 /clone="098B06"
 /clone_lib="Arabidopsis thaliana T-DNA insertion lines"
 misc_feature
 1..26
 /note="T-DNA flanking sequence
 left border"

BASE COUNT 13 a 3 c 4 g 6 t
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Query Match 100.0%; Score 10; DB 8; Length 26;
 Best Local Similarity 100.0%; Pred. No. 1.3e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
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 Db 21 CTTCTCTTTT 12

RESULT 49
 AR066258/c
 LOCUS AR066258 27 bp DNA linear PAT 29-SEP-1999
 DEFINITION Sequence 23 from patent US 5849900.
 ACCESSION AR066258
 VERSION AR066258.1 GI:5996474
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 27)
 AUTHORS Moelling,K.
 TITLE Inhibition of viruses by antisense oligomers capable of binding to polypurine rich tract of single-stranded RNA or RNA-DNA hybrids
 JOURNAL Patent: US 5849900-A 23 15-DEC-1998;
 FEATURES Location/Qualifiers
 source
 1..27
 /organism="unknown"
 BASE COUNT 7 a 8 c 8 g 4 t
 ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 27;
 Best Local Similarity 100.0%; Pred. No. 1.3e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
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 Db 22 CTTCTCTTTT 13

RESULT 50
 AR191561/c
 LOCUS AR191561 27 bp DNA linear PAT 20-APR-2002
 DEFINITION Sequence 7049 from patent US 6346398.
 ACCESSION AR191561

VERSION AR191561.1 GI:20237526
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 27)
 AUTHORS Pavco,P., McSwiggen,J., Stinchcomb,D. and Escobedo,J.
 TITLE Method and reagent for the treatment of diseases or conditions related to levels of vascular endothelial growth factor receptor
 JOURNAL Patent: US 6346398-A 7049 12-FEB-2002;
 FEATURES Location/Qualifiers
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 /organism="unknown"
 BASE COUNT 12 a 1 c 9 g 4 t 1 others
 ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 27;
 Best Local Similarity 100.0%; Pred. No. 1.3e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
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 Db 27 CTTCTCTTTT 18

RESULT 51
 AX115839/c
 LOCUS AX115839 27 bp DNA linear PAT 11-MAY-2001
 DEFINITION Sequence 962 from Patent WO0129262.
 ACCESSION AX115839
 VERSION AX115839.1 GI:14032781
 KEYWORDS
 SOURCE synthetic construct
 ORGANISM synthetic construct
 REFERENCE 1
 AUTHORS Picoult-Newburg,L. and Pohl,M.
 TITLE Genotyping reagents, kits and methods of use thereof
 JOURNAL Patent: WO 0129262-A 962 26-APR-2001;
 ORCHID Biosciences, Inc. (US)
 FEATURES Location/Qualifiers
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 /organism="synthetic construct"
 /mol_type="genomic DNA"
 /db_xref="taxon:32630"
 /note="Primer"
 BASE COUNT 13 a 5 c 4 g 5 t
 ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 27;
 Best Local Similarity 100.0%; Pred. No. 1.3e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||
 Db 21 CTTCTCTTTT 12

RESULT 52
 AX040823
 LOCUS AX040823 28 bp DNA linear PAT 23-NOV-2000
 DEFINITION Sequence 16 from Patent WO0064930.
 ACCESSION AX040823
 VERSION AX040823.1 GI:11340462
 KEYWORDS
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 REFERENCE 1
 AUTHORS Jay,G.D.
 TITLE Tribonectins
 JOURNAL Patent: WO 0064930-A 16 02-NOV-2000;

FEATURES
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RHODE ISLAND HOSPITAL (US)
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/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606" 10 t
BASE COUNT 5 a 7 c 6 g 10 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 28;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
|||||
DB 12 CTTCTCTTTT 21
RESULT 53
AR261655 29 bp DNA PAT 29-JAN-2003
LOCUS
DEFINITION Sequence 133 from patent US 6322976.
ACCESSION AR261655
VERSION AR261655.1 GI:28072733
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 29)
AUTHORS Aitman,T.J., Scott,J. and Stanton,L.W.
TITLE Compositions and methods of disease diagnosis and therapy
JOURNAL Patent: US 6322976-A 133 27-NOV-2001;
FEATURES Location/Qualifiers
source 1. .29
/organism="unknown"
BASE COUNT 1 a 5 c 10 g 13 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 29;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
|||||
DB 8 CTTCTCTTTT 17
RESULT 54
AR261654 30 bp DNA PAT 29-JAN-2003
LOCUS
DEFINITION Sequence 132 from patent US 6322976.
ACCESSION AR261654
VERSION AR261654.1 GI:28072732
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 30)
AUTHORS Aitman,T.J., Scott,J. and Stanton,L.W.
TITLE Compositions and methods of disease diagnosis and therapy
JOURNAL Patent: US 6322976-A 132 27-NOV-2001;
FEATURES Location/Qualifiers
source 1. .30
/organism="unknown"
BASE COUNT 1 a 5 c 10 g 14 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
|||||
DB 1 CTTCTCTTTT 17
RESULT 55
AR261656 30 bp DNA PAT 29-JAN-2003
LOCUS
DEFINITION Sequence 134 from patent US 6322976.
ACCESSION AR261656
VERSION AR261656.1 GI:28072734
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 30)
AUTHORS Aitman,T.J., Scott,J. and Stanton,L.W.
TITLE Compositions and methods of disease diagnosis and therapy
JOURNAL Patent: US 6322976-A 134 27-NOV-2001;
FEATURES Location/Qualifiers
source 1. .30
/organism="unknown"
BASE COUNT 1 a 5 c 10 g 14 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
|||||
DB 8 CTTCTCTTTT 17
RESULT 56
AR261657 30 bp DNA PAT 29-JAN-2003
LOCUS
DEFINITION Sequence 135 from patent US 6322976.
ACCESSION AR261657
VERSION AR261657.1 GI:28072735
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 30)
AUTHORS Aitman,T.J., Scott,J. and Stanton,L.W.
TITLE Compositions and methods of disease diagnosis and therapy
JOURNAL Patent: US 6322976-A 135 27-NOV-2001;
FEATURES Location/Qualifiers
source 1. .30
/organism="unknown"
BASE COUNT 1 a 5 c 9 g 15 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
|||||
DB 8 CTTCTCTTTT 17
RESULT 57
AR120483/c 33 bp DNA PAT 16-MAY-2001
LOCUS
DEFINITION Sequence 359 from patent US 6159469.
ACCESSION AR120483
VERSION AR120483.1 GI:14104059
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 33)
AUTHORS Choi,G.H., Kunsch,C.A., Barash,S.C., Dillon,P.J., Dougherty,B.,

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Fannon,M.R. and Rosen,C.A.
TITLE      Streptococcus pneumoniae antigens and vaccines
JOURNAL    Patent: US 6159469-A 359 12-DEC-2000;
FEATURES   Location/Qualifiers
            source          1..33
            /organism="unknown"
BASE COUNT      14 a      5 c      9 g      5 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 33;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      23 CTTCTCTTTT 14

RESULT 58
BD063492/c
LOCUS      BD063492      33 bp      DNA      linear      PAT 27-AUG-2002
DEFINITION Streptococcus pneumoniae antigens and vaccines.
ACCESSION  BD063492
VERSION     BD063492.1 GI:22609095
KEYWORDS   JP 2001505415-A/246.
SOURCE     unidentified
ORGANISM   unclassified.
REFERENCE  1 (bases 1 to 33)
AUTHORS    Kunsch,C.A., Choi,G.H., Johnson,S.L. and Hromockyj,A.
TITLE      Streptococcus pneumoniae antigens and vaccines
JOURNAL    Patent: JP 2001505415-A 246 24-APR-2001;
            HUMAN GENOME SCIENCES INC
COMMENT     PN JP 2001505415-A/246
            PD 24-APR-2001
            PF 30-OCT-1997 JP 1998520667
            PR 31-OCT-1996 US 60/029960
            PI CHARLES A KUNSCH,GIL H CHOI,SYDNOR L JOHNSON,ALEX HROMOCKYJ PC
               C12N15/31,C12N5/18,C12N1/21,C07K14/315,C12Q1/68,A61K39/09, PC
               G01N33/569,
               CC G01N33/68
               CC Strandedness: Double;
               CC Topology: Linear;
FEATURES   Location/Qualifiers
            FH Key          Location/Qualifiers
            source          1..33
            /organism="unidentified"
            /mol_type="genomic DNA"
            /db_xref="taxon:32644"
BASE COUNT      14 a      5 c      9 g      5 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 33;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      23 CTTCTCTTTT 14

RESULT 59
E36416
LOCUS      E36416      33 bp      DNA      linear      PAT 18-JUN-2001
DEFINITION dna G.
ACCESSION  E36416
VERSION     E36416.1 GI:13022642
KEYWORDS   JP 1999239489-A/3.
SOURCE     Streptococcus pneumoniae
ORGANISM   Bacteria; Firmicutes; Lactobacillales; Streptococcaceae;
            Streptococcus.
REFERENCE  1 (bases 1 to 33)

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Earl,W.M., Deborah,D.J., Ming,H., Richard,R.W. and Ana,R.R.
TITLE      dna G
JOURNAL    Patent: JP 1999239489-A 3 07-SEP-1999;
            SMITHKLINE BEECHAM CORP
COMMENT     OS Streptococcus pneumoniae
            PN JP 1999239489-A/3
            PD 07-SEP-1999
            PF 21-OCT-1998 JP 1998338366
            PR 21-OCT-1997 US 60/070912
            PI EARL WILLIAM MEI,DEBORAH D JAWASUKI,MING HWANG, PI RICHARD
               ROIDO WOREN,
               PI ANA RISA RENOX
               PC C12N15/09 A61K31/00,A61K31/70,A61K38/00,A61K39/00,A61K48/00,
               PC C07K14/315,
               PC C07K16/12,C12P21/02,G01N33/53//C12P21/08,C12N15/00,A61K37/02
               CC
               FH Key          Location/Qualifiers
               FT source      1..33
               /organism="Streptococcus pneumoniae".
FEATURES   Location/Qualifiers
            source          1..33
            /organism="Streptococcus pneumoniae"
            /mol_type="genomic DNA"
            /db_xref="taxon:1313"
BASE COUNT      8 a      7 c      4 g      14 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 33;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
            |||||
Db      11 CTTCTCTTTT 20

RESULT 60
AX218433/c
LOCUS      AX218433      38 bp      mRNA      linear      PAT 07-SEP-2001
DEFINITION Sequence 3875 from Patent WO0159103.
ACCESSION  AX218433
VERSION     AX218433.1 GI:15546157
KEYWORDS   .
SOURCE     synthetic construct
            ORGANISM       synthetic construct
                           artificial sequences.
REFERENCE  1
            BLATT,L., McSwiggen,J. and Chowrira,B.M.
            Method and reagent for the modulation and diagnosis of cd20 and
            nogo gene expression
            Patent: WO 0159103-A 3875 16-AUG-2001;
            RIBOZYME PHARMACEUTICALS, INC. (US) ; Blatt, Lawrence (US) ;
            McSwiggen, James (US) ; Chowrira, Bharat M. (US)
JOURNAL    Location/Qualifiers
FEATURES   Location/Qualifiers
            source          1..38
            /organism="synthetic construct"
            /mol_type="mRNA"
            /db_xref="taxon:32630"
            /note="Nucleic Acid"
BASE COUNT      14 a      6 c      14 g      4 t
ORIGIN

Query Match      100.0%; Score 10; DB 6; Length 38;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
            |||||
Db      38 CTTCTCTTTT 29

RESULT 61
AX220343

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LOCUS AX220343 38 bp mRNA linear PAT 07-SEP-2001
DEFINITION Sequence 5785 from Patent WO0159103.
ACCESSION AX220343
VERSION AX220343.1 GI:15548067
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1
AUTHORS Blatt, L., McSwiggen, J. and Chowrira, B.M.
TITLE Method and reagent for the modulation and diagnosis of cd20 and
nogo gene expression
JOURNAL Patent: WO 0159103-A 5785 16-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US); Blatt, Lawrence (US);
McSwiggen, James (US); Chowrira, Bharat M. (US)
FEATURES
source
1..38
Location/Qualifiers
/organism="synthetic construct"
/mol_type="mRNA"
/db_xref="taxon:32630"
/note="Nucleic Acid"
BASE COUNT 9 a 10 c 9 g 10 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 38;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 29 CTTCTCTTTT 38
RESULT 62
AX273375/c
LOCUS AX273375 38 bp mRNA linear PAT 29-OCT-2001
DEFINITION Sequence 944 from Patent WO0162911.
ACCESSION AX273375
VERSION AX273375.1 GI:16546112
KEYWORDS
SOURCE synthetic construct
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1
AUTHORS Jarvis, T., von Carlowitz, I., McSwiggen, J.A., Hamblin, P.A. and
Ellis, J.H.
TITLE Method and reagent for the inhibition of grid
JOURNAL Patent: WO 0162911-A 944 30-AUG-2001;
RIBOZYME PHARMACEUTICALS, INC. (US); GLAXO GROUP LIMITED (GB)
FEATURES
source
1..38
Location/Qualifiers
/organism="synthetic construct"
/mol_type="mRNA"
/db_xref="taxon:32630"
/note="Enzymatic Nucleic Acid"
BASE COUNT 11 a 6 c 13 g 8 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 38;
Best Local Similarity 100.0%; Pred. No. 1.3e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 38 CTTCTCTTTT 29
RESULT 63
BD182426
LOCUS BD182426 40 bp DNA linear PAT 15-MAY-2003
DEFINITION Human artificial chromosomes comprising human antibody light chain
lambda gene, and non-human animals retaining human artificial
chromosome transmittable to progeny.

ACCESSION BD182426
VERSION BD182426.1 GI:30793344
KEYWORDS WO 02092812-A/1.
SOURCE synthetic construct
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 40)
AUTHORS Kuroiwa, Y., Tomizuka, K., Yoshida, H. and Ishida, I.
TITLE Human artificial chromosomes comprising human antibody light chain
lambda gene, and non-human animals retaining human artificial
chromosome transmittable to progeny
JOURNAL Patent: WO 02092812-A 1 21-NOV-2002;
KIRIN BREWERY CO LTD, YOSHIMI KUROIWA, KAZUMA TOMIZUKA, HITOSHI
YOSHIDA, ISAO ISHIDA
COMMENT OS Artificial Sequence
PN WO 02092812-A/1
PD 21-NOV-2002
PF 10-MAY-2002 WO 2002JP004587
PR 11-MAY-2001 JP 01P 142371
PI YOSHIMI KUROIWA, KAZUMA TOMIZUKA, HITOSHI YOSHIDA, ISAO ISHIDA PC
CI2N15/09,A01K67/027,C07K16/00,C12P21/08
CC Description of Artificial Sequence:primer
FH Key Location/Qualifiers
FT source 1..40
Location/Qualifiers
/organism="Artificial Sequence".
FEATURES
source
1..40
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
BASE COUNT 5 a 14 c 6 g 15 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 40;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 21 CTTCTCTTTT 30
RESULT 64
I12092
LOCUS I12092 42 bp DNA linear PAT 26-JUL-1995
DEFINITION Sequence 33 from patent US 5420027.
ACCESSION I12092
VERSION I12092.1 GI:909590
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 42)
AUTHORS Fisher, C.W., Barnes, H.J. and Estabrook, R.W.
TITLE Methods and compositions for the expression of biologically active
fusion proteins comprising a eukaryotic cytochrome P450 fused to a
reductase in bacteria
JOURNAL Patent: US 5420027-A 33 30-MAY-1995;
FEATURES
source
1..42
Location/Qualifiers
/organism="unknown"
BASE COUNT 5 a 9 c 7 g 21 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 42;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 34

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RESULT 65
AX484623
LOCUS
DEFINITION Sequence 1923 from Patent WO02053728.
ACCESSION AX484623
VERSION AX484623.1 GI:22318975
KEYWORDS
SOURCE
ORGANISM Candida albicans
Candida albicans
Bukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
Saccharomycetales; mitosporic Saccharomycetales; Candida.
REFERENCE
AUTHORS Roemer, T., Jiang, B., Boone, C., Bussey, H. and Ohlsen, K.L.
TITLE Gene disruption methodologies for drug target discovery
JOURNAL Patent: WO 02053728-A 1923 11-JUL-2002;
Elitra Pharmaceuticals, Inc. (US)
FEATURES
source
Location/Qualifiers
1..43
/organism="Candida albicans"
/mol_type="genomic DNA"
/db_xref="taxon:5476" 29 t
BASE COUNT 7 a 5 c 2 g 29 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 43;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 12 CTTCTCTTTT 21

RESULT 66
AX061875/c
LOCUS
DEFINITION Sequence 8 from Patent WO0078978.
ACCESSION AX061875
VERSION AX061875.1 GI:12539921
KEYWORDS
SOURCE
ORGANISM synthetic construct
synthetic construct
artificial sequences.
REFERENCE
AUTHORS Miller, B.G., Sloan, J.S., Raymond, C.K. and Vanaja, E.
TITLE Pichia methanolica glycerolaldehyde-3-phosphate dehydrogenase 1
JOURNAL Patent: WO 0078978-A 8 28-DEC-2000;
ZymoGenetics, Inc. (US) ; Miller, Brady G. (US) ; Sloan, James S.
(US)
FEATURES
source
Location/Qualifiers
1..45
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
/Note="oligonucleotide primer ZC12.565"
BASE COUNT 23 a 9 c 8 g 5 t
ORIGIN
Query Match 100.0%; Score 10; DB 6; Length 45;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 16 CTTCTCTTTT 7

RESULT 67
ATH528602/c
LOCUS
DEFINITION Arabidopsis thaliana T-DNA flanking sequence, left border, clone
168E06.

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ACCESSION AJ528602
VERSION AJ528602.1 GI:26796862
KEYWORDS left border; T-DNA flanking sequence.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Arabidopsi
Bukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi
1
Brunaud, V., Balzergue, S., Dubreucq, B., Aubourg, S., Samson, F.,
Chauvin, S., Bechtold, N., Cruaud, C., DeRose, R., Pelletier, G.,
Lepiniec, L., Caboche, M. and Lecharny, A.
T-DNA integration into the Arabidopsis genome depends on sequences
of pre-insertion sites
EMBO Rep. 3 (12), 1152-1157 (2002)
MEDLINE 22363535
PUBMED 1246565
REFERENCE 2 (bases 1 to 45)
AUTHORS Balzergue, S.
TITLE Direct Submission
JOURNAL Submitted (21-NOV-2002) Balzergue S., UMRGV, INRA/CNRS, 2 rue
Gaston Cremieux, 91057 Evry cedex, FRANCE
COMMENT PCR was performed on DNA from transformants of Arabidopsis thaliana
plants from INRA (Versailles). The DNA fragment(s) resulting from
the PCR were directly sequenced from the left or the right border
to determine the genomic sequence flanking the insertion. T-DNA
derived sequences were removed. Information to order the
corresponding mutant line and a link to a database providing a
graphical display of the insertion site are available at
http://dbsgap.versailles.inra.fr/publiclines/. This sequence has
been generated in the framework of the French plant genomics
program 'Genoplante' (http://www.genoplante.com and
http://genoplante-info.infobiogen.fr).
FEATURES
source
1..45
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/cultivar="Wassillewskija"
/db_xref="taxon:3702"
/clone="168E06"
/Note="T-DNA flanking sequence"
misc_feature 1..45
/Note="T-DNA flanking sequence"
left border"
BASE COUNT 16 a 8 c 9 g 12 t
ORIGIN
Query Match 100.0%; Score 10; DB 8; Length 45;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 10 CTTCTCTTTT 1

RESULT 68
AR284713
LOCUS
DEFINITION Sequence 765 from patent US 6528260.
ACCESSION AR284713
VERSION AR284713.1 GI:29721617
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
Unclassified.
REFERENCE 1 (bases 1 to 47)
AUTHORS Blumenfeld, M., Chumakov, I., Bougueleret, L. and Cohen, A.
TITLE Biallelic markers related to genes involved in drug metabolism
JOURNAL Patent: US 6528260-A 765 04-MAR-2003;
FEATURES
source
Location/Qualifiers
1..47
/organism="unknown"

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BASE COUNT 6 a 11 c 8 g 21 t 1 others
 ORIGIN
 Query Match 100.0%; Score 10; DB 6; Length 47;
 Best Local Similarity 100.0%; Pred. No. 1.2e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 1 CTTCTCTTTT 10
 RESULT 69
 AR288787/c
 LOCUS AR288787 47 bp DNA PAT 12-JUN-2003
 DEFINITION Sequence 522 from patent US 6537751.
 ACCESSION AR288787
 VERSION AR288787.1 GI:31676071
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 47)
 AUTHORS Cohen, D., Chumakov, I. and Blumenfeld, M.
 TITLE Biallelic markers for use in constructing a high density
 disequilibrium map of the human genome
 JOURNAL Patent: US 6537751-A 522 25-MAR-2003;
 FEATURES Location/Qualifiers
 source 1..47
 BASE COUNT 16 a 4 c 18 g 8 t 1 others
 ORIGIN
 Query Match 100.0%; Score 10; DB 6; Length 47;
 Best Local Similarity 100.0%; Pred. No. 1.2e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 15 CTTCTCTTTT 6
 RESULT 70
 AR289547
 LOCUS AR289547 47 bp DNA PAT 12-JUN-2003
 DEFINITION Sequence 1282 from patent US 6537751.
 ACCESSION AR289547
 VERSION AR289547.1 GI:31676831
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 47)
 AUTHORS Cohen, D., Chumakov, I. and Blumenfeld, M.
 TITLE Biallelic markers for use in constructing a high density
 disequilibrium map of the human genome
 JOURNAL Patent: US 6537751-A 1282 25-MAR-2003;
 FEATURES Location/Qualifiers
 source 1..47
 BASE COUNT 9 a 12 c 10 g 15 t 1 others
 ORIGIN
 Query Match 100.0%; Score 10; DB 6; Length 47;
 Best Local Similarity 100.0%; Pred. No. 1.2e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 13 CTTCTCTTTT 22
 RESULT 71

AR289811
 LOCUS AR289811 47 bp DNA PAT 12-JUN-2003
 DEFINITION Sequence 1546 from patent US 6537751.
 ACCESSION AR289811
 VERSION AR289811.1 GI:31677095
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 47)
 AUTHORS Cohen, D., Chumakov, I. and Blumenfeld, M.
 TITLE Biallelic markers for use in constructing a high density
 disequilibrium map of the human genome
 JOURNAL Patent: US 6537751-A 1546 25-MAR-2003;
 FEATURES Location/Qualifiers
 source 1..47
 BASE COUNT 5 a 14 c 24 t 2 others
 ORIGIN
 Query Match 100.0%; Score 10; DB 6; Length 47;
 Best Local Similarity 100.0%; Pred. No. 1.2e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 9 CTTCTCTTTT 18
 RESULT 72
 AR290737/c
 LOCUS AR290737 47 bp DNA PAT 12-JUN-2003
 DEFINITION Sequence 2472 from patent US 6537751.
 ACCESSION AR290737
 VERSION AR290737.1 GI:31678021
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 47)
 AUTHORS Cohen, D., Chumakov, I. and Blumenfeld, M.
 TITLE Biallelic markers for use in constructing a high density
 disequilibrium map of the human genome
 JOURNAL Patent: US 6537751-A 2472 25-MAR-2003;
 FEATURES Location/Qualifiers
 source 1..47
 BASE COUNT 24 a 6 c 7 g 9 t 1 others
 ORIGIN
 Query Match 100.0%; Score 10; DB 6; Length 47;
 Best Local Similarity 100.0%; Pred. No. 1.2e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 45 CTTCTCTTTT 36
 RESULT 73
 AR291549/c
 LOCUS AR291549 47 bp DNA PAT 12-JUN-2003
 DEFINITION Sequence 3284 from patent US 6537751.
 ACCESSION AR291549
 VERSION AR291549.1 GI:31678833
 KEYWORDS
 SOURCE Unknown.
 ORGANISM Unknown.
 REFERENCE 1 (bases 1 to 47)
 AUTHORS Cohen, D., Chumakov, I. and Blumenfeld, M.
 TITLE Biallelic markers for use in constructing a high density
 disequilibrium map of the human genome

JOURNAL Patent: US 6537751-A 3284 25-MAR-2003;
FEATURES
source
1. .47
Location/Qualifiers
/organism="unknown"
BASE COUNT 19 a 4 c 12 g 11 t 1 others
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 41 CTTCTCTTTT 32

RESULT 74
AR139658/c
LOCUS AR139658 50 bp DNA linear PAT 16-JUN-2001
DEFINITION Sequence 37 from patent US 6207389.
ACCESSION AR139658
VERSION AR139658.1 GI:14482154
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 50)
AUTHORS Dosch,H.Michael.
TITLE Methods of controlling T lymphocyte mediated immune responses
JOURNAL Patent: US 6207389-A 37 27-MAR-2001;
FEATURES
source
1. .50
/organism="unknown"
BASE COUNT 20 a 8 c 16 g 6 t
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 50;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 42 CTTCTCTTTT 33

RESULT 75
AX158156
LOCUS AX158156 50 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 1484 from Patent WO0140521.
ACCESSION AX158156
VERSION AX158156.1 GI:14539487
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1
AUTHORS Shimkets,R.A. and Leach,M.
TITLE Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
JOURNAL Patent: WO 0140521-A 1484 07-JUN-2001;
Curagen Corporation (US)
FEATURES
source
1. .50
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
misc_feature 25..26
/note="Nucleotide deleted between bases 25 and 26
Accession number cg2968983"
misc_feature 26
/note="2 of 2 allelic variants (1483 is other entry)"
BASE COUNT 4 a 9 c 2 g 35 t

ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 50;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 26 CTTCTCTTTT 35

RESULT 76
AX118141
LOCUS AX118141 51 bp DNA linear PAT 11-MAY-2001
DEFINITION Sequence 3264 from Patent WO0129262.
ACCESSION AX118141
VERSION AX118141.1 GI:14035092
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1
AUTHORS Picoult-Newburg,L. and Pohl,M.
TITLE Genotyping reagents, kits and methods of use thereof
JOURNAL Patent: WO 0129262-A 3264 26-APR-2001;
Orchid Biosciences, Inc. (US)
FEATURES
source
1. .51
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
BASE COUNT 8 a 13 c 3 g 27 t
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 51;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 11 CTTCTCTTTT 20

RESULT 77
AX158155
LOCUS AX158155 51 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 1483 from Patent WO0140521.
ACCESSION AX158155
VERSION AX158155.1 GI:14539486
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1
AUTHORS Shimkets,R.A. and Leach,M.
TITLE Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
JOURNAL Patent: WO 0140521-A 1483 07-JUN-2001;
Curagen Corporation (US)
FEATURES
source
1. .51
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
misc_feature 26
/note="1 of 2 allelic variants (1484 is other entry)"
BASE COUNT 4 a 9 c 2 g 36 t
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 51;

Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 27 CTTCTCTTTT 36

RESULT 78
AX160381
LOCUS AX160381 51 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 3709 from Patent WO0140521.
ACCESSION AX160381
VERSION AX160381.1 GI:14541712
KEYWORDS Homo sapiens (human)
SOURCE
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1
AUTHORS Shimkets, R.A. and Leach, M.
TITLE Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
JOURNAL Patent: WO 0140521-A 3709 07-JUN-2001;
Curagen Corporation (US)
FEATURES
source Location/Qualifiers
1..51
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
misc_feature 26
/note="1 of 2 allelic variants (3710 is other entry)
Accession number CG43917418"
BASE COUNT 7 a 15 c 10 g 19 t
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 51;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 40 CTTCTCTTTT 49

RESULT 81
AX165562
LOCUS AX165562 51 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 757 from Patent WO0138586.
ACCESSION AX165562
VERSION AX165562.1 GI:14546391
KEYWORDS Homo sapiens (human)
SOURCE
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1
AUTHORS Shimkets, R.A. and Leach, M.
TITLE Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
JOURNAL Patent: WO 0138586-A 757 31-MAY-2001;
Curagen Corporation (US)
FEATURES
source Location/Qualifiers
1..51
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
misc_feature 26
/note="2 of 2 allelic variants (3709 is other entry)
Accession number CG43917418"
BASE COUNT 7 a 16 c 10 g 18 t
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 51;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 40 CTTCTCTTTT 49

RESULT 80
AX160383
LOCUS AX160383 51 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 3711 from Patent WO0140521.
ACCESSION AX160383
VERSION AX160383.1 GI:14541714
KEYWORDS Homo sapiens (human)
SOURCE
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1
AUTHORS Shimkets, R.A. and Leach, M.
TITLE Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
JOURNAL Patent: WO 0140521-A 3711 07-JUN-2001;
Curagen Corporation (US)
FEATURES
source Location/Qualifiers
1..51
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
misc_feature 26
/note="1 of 2 allelic variants (3712 is other entry)
Accession number CG43917418"
BASE COUNT 6 a 12 c 10 g 23 t
ORIGIN

Query Match 100.0%; Score 10; DB 6; Length 51;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 25 CTTCTCTTTT 34

RESULT 81
AX165562
LOCUS AX165562 51 bp DNA linear PAT 22-JUN-2001
DEFINITION Sequence 757 from Patent WO0138586.
ACCESSION AX165562
VERSION AX165562.1 GI:14546391
KEYWORDS Homo sapiens (human)
SOURCE
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1
AUTHORS Shimkets, R.A. and Leach, M.
TITLE Nucleic acids containing single nucleotide polymorphisms and methods of use thereof
JOURNAL Patent: WO 0138586-A 757 31-MAY-2001;
Curagen Corporation (US)
FEATURES
source Location/Qualifiers
1..51
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
variation 26
/note="single nucleotide polymorphism
Accession number CG43917191"
BASE COUNT 12 a 12 c 7 g 20 t
ORIGIN

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Query Match      100.0%; Score 10; DB 6; Length 51;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 2 CTTCTCTTTT 11

RESULT 82
ATH521150
LOCUS      51 bp      DNA      linear      PLN 29-MAR-2003
DEFINITION Arabidopsis thaliana T-DNA flanking sequence, left border, clone
            260A09.
ACCESSION  AJ521150
VERSION     AJ521150.1 GI:26789386
KEYWORDS   left border; T-DNA flanking sequence.
SOURCE     Arabidopsis thaliana (thale cress)
ORGANISM   Arabidopsis thaliana
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
            rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
1
REFERENCE   1
AUTHORS    Brunaud,V., Balzergue,S., Dubreucq,B., Aubourg,S., Samson,F.,
            Chauvin,S., Reichtold,N., Cruaud,C., DeRose,R., Pelletier,G.,
            Lepiniec,L., Caboche,M. and Lecharny,A.
TITLE      T-DNA integration into the Arabidopsis genome depends on sequences
            of pre-insertion sites
JOURNAL    EMBO Rep. 3 (12), 1152-1157 (2002)
MEDLINE    22363535
PUBMED     12446565
REFERENCE   2 (bases 1 to 51)
AUTHORS    Balzergue,S.
TITLE      Direct Submission
JOURNAL    Submitted (21-NOV-2002) Balzergue S., UMRGV, INRA/CNRS, 2 rue
            Gaston Cremieux, 91057 Evry cedex, FRANCE
COMMENT    PCR was performed on DNA from transformants of Arabidopsis thaliana
            plants from INRA (Versailles). The DNA fragment(s) resulting from
            the PCR were directly sequenced from the left or the right border
            to determine the genomic sequence flanking the insertion. T-DNA
            derived sequences were removed. Information to order the
            corresponding mutant line and a link to a database providing a
            graphical display of the insertion site are available at
            http://dbgap.versailles.inra.fr/publiclines/. This sequence has
            been generated in the framework of the French plant genomics
            program 'Genoplante' (http://www.genoplante.com and
            http://genoplante-info.infobiogen.fr).
            Location/Qualifiers
            1..51
            /organism="Arabidopsis thaliana"
            /mol_type="genomic DNA"
            /cultivar="Wassilewskija"
            /db_xref="taxon:3702"
            /clone="260A09"
            /clone_lib="Arabidopsis thaliana T-DNA insertion lines"
            misc_feature
            1..51
            /note="T-DNA flanking sequence
            left border"
BASE COUNT      15 a      9 c      2 g      25 t
ORIGIN
Query Match      100.0%; Score 10; DB 8; Length 51;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 4 CTTCTCTTTT 13

RESULT 83
AR098682/c
LOCUS      53 bp      DNA      linear      PAT 14-FEB-2001
DEFINITION Sequence 40 from patent US 6077668.
ACCESSION  AR098682
VERSION     AR098682.1 GI:12808448
KEYWORDS   Unknown.
SOURCE     Unknown.
ORGANISM   Unclassified.
            1 (bases 1 to 53)
AUTHORS    Kool,E.T.
TITLE      Highly sensitive multimeric nucleic acid probes
JOURNAL    Patent: US 6077668-A 40 20-JUN-2000;
            Location/Qualifiers
            1..53
            /organism="unknown"
BASE COUNT      20 a      10 c      15 g      8 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 25 CTTCTCTTTT 16

RESULT 84
AR098683
LOCUS      53 bp      DNA      linear      PAT 14-FEB-2001
DEFINITION Sequence 41 from patent US 6077668.
ACCESSION  AR098683
VERSION     AR098683.1 GI:12808449
KEYWORDS   Unknown.
SOURCE     Unknown.
ORGANISM   Unclassified.
            1 (bases 1 to 53)
AUTHORS    Kool,E.T.
TITLE      Highly sensitive multimeric nucleic acid probes
JOURNAL    Patent: US 6077668-A 41 20-JUN-2000;
            Location/Qualifiers
            1..53
            /organism="unknown"
BASE COUNT      8 a      15 c      10 g      20 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 38 CTTCTCTTTT 47

RESULT 85
AR204756/c
LOCUS      53 bp      DNA      linear      PAT 20-JUN-2002
DEFINITION Sequence 40 from patent US 638802.
ACCESSION  AR204756
VERSION     AR204756.1 GI:21502164
KEYWORDS   Unknown.
SOURCE     Unknown.
ORGANISM   Unclassified.
            1 (bases 1 to 53)
AUTHORS    Kool,E.T.
TITLE      Circular DNA vectors for synthesis of RNA and DNA
JOURNAL    Patent: US 638802-A 40 09-APR-2002;
            Location/Qualifiers
            1..53
            /organism="unknown"
            source

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BASE COUNT      20 a      10 c      15 g      8 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      25 CTTCTCTTTT 16

RESULT 86
AR204757
LOCUS      AR204757      53 bp      DNA      linear      PAT 20-JUN-2002
DEFINITION Sequence 41 from patent US 6368802.
ACCESSION AR204757
VERSION AR204757.1 GI:21502165
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 53)
AUTHORS Kool,E.T.
TITLE Circular DNA vectors for synthesis of RNA and DNA
JOURNAL Patent: US 6368802-A 41 09-APR-2002;
FEATURES Location/Qualifiers
          source
            1..53
              /organism="unknown"
BASE COUNT      8 a      15 c      10 g      20 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      38 CTTCTCTTTT 47

RESULT 87
AR134108/c
LOCUS      AR134108      54 bp      DNA      linear      PAT 16-MAY-2001
DEFINITION Sequence 2533 from patent US 6194150.
ACCESSION AR134108
VERSION AR134108.1 GI:14123013
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 54)
AUTHORS Stinchcomb,D.T., Jarvis,T. and McSwiggen,J.
TITLE Nucleic acid based inhibition of CD40
JOURNAL Patent: US 6194150-A 2533 27-FEB-2001;
FEATURES Location/Qualifiers
          source
            1..54
              /organism="unknown"
BASE COUNT      20 a      12 c      13 g      9 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 54;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      13 CTTCTCTTTT 4

RESULT 88
AR134285/c
LOCUS      AR134285      54 bp      DNA      linear      PAT 16-MAY-2001

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DEFINITION Sequence 2710 from patent US 6194150.
ACCESSION AR134285
VERSION AR134285.1 GI:14123190
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 54)
AUTHORS Stinchcomb,D.T., Jarvis,T. and McSwiggen,J.
TITLE Nucleic acid based inhibition of CD40
JOURNAL Patent: US 6194150-A 2710 27-FEB-2001;
FEATURES Location/Qualifiers
          source
            1..54
              /organism="unknown"
BASE COUNT      21 a      11 c      12 g      10 t
ORIGIN
Query Match      100.0%; Score 10; DB 6; Length 54;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      13 CTTCTCTTTT 4

RESULT 89
ATH505724
LOCUS      ATH505724      57 bp      mRNA      linear      PLN 30-APR-2003
DEFINITION Arabidopsis thaliana partial mitochondrial small non-messenger RNA, clone Ath-647, 5' end incomplete.
ACCESSION AJ505724
VERSION AJ505724.1 GI:22293620
KEYWORDS small non-messenger RNA; smRNA.
SOURCE Mitochondrion Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
REFERENCE 1
AUTHORS Marker,C., Zemann,A., Terhorst,T., Kiefmann,M., Kastenmayer,J.P., Green,P., Bachelierie,J.P., Brosius,J. and Huttenhofer,A.
TITLE Experimental RNomics. Identification of 140 Candidates for Small Non-Messenger RNAs in the Plant Arabidopsis thaliana
JOURNAL Curr. Biol. 12 (23), 2002-2013 (2002)
MEDLINE 22365595
PUBMED 12477388
REFERENCE 2 (bases 1 to 57)
AUTHORS Huttenhofer,A.
TITLE Direct Submission
JOURNAL Submitted (22-JUL-2002) Huttenhofer A., University of Muenster, Institute for Experimental Pathology, Von-Esmarch-Str. 56, 48149 Muenster, GERMANY
FEATURES Location/Qualifiers
          source
            1..57
              /organism="Arabidopsis thaliana"
              /organelle="mitochondrion"
              /mol_type="mRNA"
              /db_xref="taxon:3702"
              /clone="Ath-647"
              /note="small non-messenger RNA, smRNA"
BASE COUNT      6 a      10 c      8 g      33 t
ORIGIN
Query Match      100.0%; Score 10; DB 8; Length 57;
Best Local Similarity 100.0%; Pred. No. 1.2e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      45 CTTCTCTTTT 54

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RESULT 90
ATH527067/c
LOCUS
DEFINITION
  ATH527067          58 bp      DNA      linear      PLN 29-MAR-2003
  Arabidopsis thaliana T-DNA flanking sequence, left border, clone
  131H07
ACCESSION
  AJ527067          GI:26795327
VERSION
  AJ527067.1
KEYWORDS
  left border; T-DNA flanking sequence.
SOURCE
  Arabidopsis thaliana (thale cress)
ORGANISM
  Arabidopsis thaliana
  Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
  Spermatophyta; Magnoliophyta, eudicotyledons, core eudicots;
  rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE
  1
  Brunaud,V., Balzergue,S., Dubreucq,B., Aubourg,S., Samson,F.,
  Chauvin,S., Bechtold,N., Cruaud,C., DeRose,R., Pelletier,G.,
  Lepiniec,L., Caboche,M. and Lecharny,A.
  T-DNA integration into the Arabidopsis genome depends on sequences
  of pre-insertion sites
  EMBO Rep. 3 (12), 1152-1157 (2002)
JOURNAL
  22363535
MEDLINE
  12446565
PUBMED
  12446565
REFERENCE
  2 (bases 1 to 58)
  Balzergue,S.
  Direct Submission
  Submitted (21-NOV-2002) Balzergue S., UMRGV, INRA/CNRS, 2 rue
  Gaston Cremieux, 91057 Evry cedex, FRANCE
COMMENT
  PCR was performed on DNA from transformants of Arabidopsis thaliana
  plants from INRA (Versailles). The DNA fragment (s) resulting from
  the PCR were directly sequenced from the left or the right border
  to determine the genomic sequence flanking the insertion. T-DNA
  derived sequences were removed. Information to order the
  corresponding mutant line and a link to a database providing a
  graphical display of the insertion site are available at
  http://dbsgap.versailles.inra.fr/publiclines/. This sequence has
  been generated in the framework of the French plant genomics
  program 'Genoplante' (http://www.genoplante.com and
  http://genoplante-info.infobiogen.fr).
FEATURES
  source
  1..58
  /organism="Arabidopsis thaliana"
  /mol_type="genomic DNA"
  /cultivar="Wassilewskija"
  /db_xref="taxon:3702"
  /clone="131H07"
  /clone_lib="Arabidopsis thaliana T-DNA insertion lines"
  misc_feature
  1..58
  /note="T-DNA flanking sequence
  left border"
BASE COUNT
  18 a      4 c      14 g      22 t
ORIGIN
  Query Match          100.0%; Score 10; DB 8; Length 58;
  Best Local Similarity 100.0%; Pred. No. 1.2e+05;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  Qy 1 CTTCTCTTTT 10
  Db 47 CTTCTCTTTT 38

RESULT 91
AX270701
LOCUS
DEFINITION
  Sequence 1332 from Patent WO0164876.
ACCESSION
  AX270701
VERSION
  AX270701.1
KEYWORDS
  Homo sapiens (human)
SOURCE
  Homo sapiens
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
  1
  Stefansson,H., Steinthorsdottir,V. and Gulcher,J.R.
  Human schizophrenia gene
  Patent: WO 0164876-A 1332 07-SEP-2001;
  Decode Genetics EHF. (IS)
FEATURES
  source
  1..61
  /organism="Homo sapiens"
  /mol_type="genomic DNA"
  /db_xref="taxon:9606"
BASE COUNT
  2 a      23 c      0 g      35 t      1 others
ORIGIN
  Query Match          100.0%; Score 10; DB 6; Length 61;
  Best Local Similarity 100.0%; Pred. No. 1.2e+05;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  Qy 1 CTTCTCTTTT 10
  Db 17 CTTCTCTTTT 26

RESULT 92
AX272232
LOCUS
DEFINITION
  Sequence 1332 from Patent WO0164877.
ACCESSION
  AX272232
VERSION
  AX272232.1
KEYWORDS
  Homo sapiens (human)
SOURCE
  Homo sapiens
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
  1
  Stefansson,H., Steinthorsdottir,V. and Gulcher,J.R.
  Human schizophrenia gene
  Patent: WO 0164877-A 1332 07-SEP-2001;
  Decode Genetics EHF. (IS)
FEATURES
  source
  1..61
  /organism="Homo sapiens"
  /mol_type="genomic DNA"
  /db_xref="taxon:9606"
BASE COUNT
  2 a      23 c      0 g      35 t      1 others
ORIGIN
  Query Match          100.0%; Score 10; DB 6; Length 61;
  Best Local Similarity 100.0%; Pred. No. 1.2e+05;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  Qy 1 CTTCTCTTTT 10
  Db 17 CTTCTCTTTT 26

RESULT 93
AX482835
LOCUS
DEFINITION
  Sequence 135 from Patent WO02053728.
ACCESSION
  AX482835
VERSION
  AX482835.1
KEYWORDS
  Candida albicans
SOURCE
  Candida albicans
  Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
  Saccharomycetales; mitosporic Saccharomycetales; Candida.
REFERENCE
  1
  Roemer,T., Jiang,B., Boone,C., Bussey,H. and Ohlsen,K.L.
  Gene disruption methodologies for drug target discovery
  Patent: WO 02053728-A 135 11-JUL-2002;
  Elitra Pharmaceuticals, Inc. (US)
FEATURES
  source
  1..65
  /organism="Homo sapiens"
  /mol_type="genomic DNA"
  /db_xref="taxon:9606"
BASE COUNT
  2 a      23 c      0 g      35 t      1 others
ORIGIN
  Query Match          100.0%; Score 10; DB 6; Length 61;
  Best Local Similarity 100.0%; Pred. No. 1.2e+05;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

  Qy 1 CTTCTCTTTT 10
  Db 17 CTTCTCTTTT 26

RESULT 94
AX482835
LOCUS
DEFINITION
  Sequence 135 from Patent WO02053728.
ACCESSION
  AX482835
VERSION
  AX482835.1
KEYWORDS
  Candida albicans
SOURCE
  Candida albicans
  Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
  Saccharomycetales; mitosporic Saccharomycetales; Candida.
REFERENCE
  1
  Roemer,T., Jiang,B., Boone,C., Bussey,H. and Ohlsen,K.L.
  Gene disruption methodologies for drug target discovery
  Patent: WO 02053728-A 135 11-JUL-2002;
  Elitra Pharmaceuticals, Inc. (US)
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DEFINITION Sequence 152 from Patent WO202053728.
ACCESSION  AX482852
VERSION     AX482852.1 GI:22317272
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SOURCE     Candida albicans
ORGANISM   Candida albicans
REFERENCE  Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
            Saccharomycetales; mitosporic Saccharomycetales; Candida.
AUTHORS    Roemer,T., Jiang,B., Boone,C., Bussey,H. and Ohlsen,K.L.
TITLE      Gene disruption methodologies for drug target discovery
JOURNAL    Patent: WO 02053728-A 152 11-JUL-2002;
            Elitra Pharmaceuticals, Inc. (US)
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DEFINITION Sequence 2790 from Patent WO202053728.
ACCESSION  AX485490
VERSION     AX485490.1 GI:22319774
KEYWORDS   .
SOURCE     Candida albicans
ORGANISM   Candida albicans
REFERENCE  Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
            Saccharomycetales; mitosporic Saccharomycetales; Candida.
AUTHORS    Roemer,T., Jiang,B., Boone,C., Bussey,H. and Ohlsen,K.L.
TITLE      Gene disruption methodologies for drug target discovery
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DEFINITION Sequence 1 from Patent WO9900504.
ACCESSION  A81696
VERSION     A81696.1 GI:6731831
KEYWORDS   .
SOURCE     unidentified
ORGANISM   unidentified
REFERENCE  1 (bases 1 to 70)
AUTHORS    Sleep,D.
TITLE      IMPROVED PROTEIN EXPRESSION STRAINS
JOURNAL    Patent: WO 9900504-A 1 07-JAN-1999;
            DELTA BIOTECHNOLOGY LTD (GB); SLEEP DARRELL (GB)
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VERSION     AR207788.1 GI:21507632
KEYWORDS   .
SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 70)
AUTHORS    Sleep,D.
TITLE      Protein expression strains
JOURNAL    Patent: US 6379924-A 1 30-APR-2002;
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RESULT 98
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DEFINITION Sequence 5 from patent US 5659122.

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ACCESSION      I62432
VERSION        I62432.1  GI:2480380
KEYWORDS
SOURCE         Unknown.
ORGANISM       Unclassified.
REFERENCE      1 (bases 1 to 71)
AUTHORS        Austin,G.Douglas.
TITLE          Enhanced expression in plants using non-translated leader sequences
JOURNAL        Patent: US 5659122-A 5 19-AUG-1997;
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DEFINITION    Homo sapiens mitochondrial short-chain L-3-hydroxyacyl-CoA
               dehydrogenase (HADHSC) gene, nuclear gene encoding mitochondrial
               protein, 5' end of intron 3.
ACCESSION     AF026858
VERSION       AF026858.1  GI:3882441
KEYWORDS
SEGMENT
SOURCE        6 of 15
ORGANISM      Homo sapiens (human)
               Homo sapiens
REFERENCE      1 (bases 1 to 71)
AUTHORS        Vredendaal,P.J., van den Berg,I.E., Stroobants,A.K., van der
               A.D.L., Malingre,H.E. and Berger,R.
TITLE          Structural organization of the human short-chain
               L-3-hydroxyacyl-CoA dehydrogenase gene
JOURNAL        Mamm. Genome 9 (9), 763-768 (1998)
MEDLINE       98384544
PUBMED        9716664
REFERENCE      2 (bases 1 to 71)
AUTHORS        Vredendaal,P.J.C.M., van den Berg,I.E.T., Stroobants,A.K., van der
               A.D.L., Malingre,H.E.M. and Berger,R.
TITLE          Direct Submission
JOURNAL        Submitted (25-SEP-1997) Department of Metabolic Diseases,
               Wilhelmina Children's Hospital, Nieuwe Gracht 137, Utrecht 3512 LK,
               The Netherlands
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Job time : 1553 secs

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ACCESSION     BD055583
VERSION       BD055583.1  GI:22601189
KEYWORDS      JP 2001269182-A/31829.
SOURCE        Homo sapiens (human)
ORGANISM      Homo sapiens
               Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
               Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE      1 (bases 1 to 73)
AUTHORS        Edwards,J.B.D.M., Duclair,E. and Jordan,J.Y.
TITLE          Sequence tag and encoded human protein
JOURNAL        Patent: JP 2001269182-A 31829 02-OCT-2001;
               GENSET
COMMENT        OS Homo sapiens (human)
               PN JP 2001269182-A/31829
               PD 02-OCT-2001
               PF 24-FEB-2000 JP 2000118773
               PR 26-FEB-1999 US 60/122487
               PI JEAN BAPTISTE DUMAS MILNE EDWARDS,EIMERIC DUCLAIR,JEAN YVES
               PI JORDAN
               PC C12N15/09,C07K14/435,C07K16/18,C12N1/15,C12N1/19,C12N1/21, PC
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Search completed: October 28, 2003, 17:43:55
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

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C 118	10	100.0	42	13	AAQ27383	Bovine 17-alpha-hy	C 191	10	100.0	87	22	AAH86210	Human single nucle
C 119	10	100.0	42	15	AAQ55121	Human liver P450 2	C 192	10	100.0	87	22	AAH86210	Human single nucle
C 120	10	100.0	42	20	AAH86210	Human single nucle	C 193	10	100.0	87	22	AAH86210	Human single nucle
C 121	10	100.0	43	24	ABZ27976	Candida essential	C 194	10	100.0	88	22	AAH86210	Human single nucle
C 122	10	100.0	45	22	AAH86210	Human single nucle	C 195	10	100.0	89	23	AAH86210	Human single nucle
C 123	10	100.0	47	20	AAH86210	Human single nucle	C 196	10	100.0	89	23	AAH86210	Human single nucle
C 124	10	100.0	47	21	AAH86210	Human single nucle	C 197	10	100.0	90	24	AAH86210	Human single nucle
C 125	10	100.0	47	21	AAH86210	Human single nucle	C 198	10	100.0	93	22	AAH86210	Human single nucle
C 126	10	100.0	47	21	AAH86210	Human single nucle	C 199	10	100.0	93	22	AAH86210	Human single nucle
C 127	10	100.0	47	21	AAH86210	Human single nucle	C 200	10	100.0	93	22	AAH86210	Human single nucle
C 128	10	100.0	47	21	AAH86210	Human single nucle	C 201	10	100.0	93	22	AAH86210	Human single nucle
C 129	10	100.0	48	21	AAH86210	Human single nucle	C 202	10	100.0	93	22	AAH86210	Human single nucle
C 130	10	100.0	50	21	AAH86210	Human single nucle	C 203	10	100.0	93	22	AAH86210	Human single nucle
C 131	10	100.0	50	21	AAH86210	Human single nucle	C 204	10	100.0	93	22	AAH86210	Human single nucle
C 132	10	100.0	50	22	AAH86210	Human single nucle	C 205	10	100.0	93	22	AAH86210	Human single nucle
C 133	10	100.0	50	22	AAH86210	Human single nucle	C 206	10	100.0	93	22	AAH86210	Human single nucle
C 134	10	100.0	50	24	ABZ02704	Human leukocyte ge	C 207	10	100.0	96	22	AAH86210	Human single nucle
C 135	10	100.0	50	24	ABZ04201	Human leukocyte ge	C 208	10	100.0	96	22	AAH86210	Human single nucle
C 136	10	100.0	50	24	ABZ04416	Human leukocyte ge	C 209	10	100.0	97	22	AAH86210	Human single nucle
C 137	10	100.0	50	24	ABZ06159	Human leukocyte ge	C 210	10	100.0	97	22	AAH86210	Human single nucle
C 138	10	100.0	50	24	ABZ06269	Human leukocyte ge	C 211	10	100.0	97	22	AAH86210	Human single nucle
C 139	10	100.0	51	21	AAH86210	Human single nucle	C 212	10	100.0	97	22	AAH86210	Human single nucle
C 140	10	100.0	51	22	AAH86210	Human single nucle	C 213	10	100.0	97	22	AAH86210	Human single nucle
C 141	10	100.0	51	22	AAH86210	Human single nucle	C 214	10	100.0	97	22	AAH86210	Human single nucle
C 142	10	100.0	51	22	AAH86210	Human single nucle	C 215	10	100.0	97	22	AAH86210	Human single nucle
C 143	10	100.0	51	22	AAH86210	Human single nucle	C 216	10	100.0	99	21	AAH86210	Human single nucle
C 144	10	100.0	51	22	AAH86210	Human single nucle	C 217	10	100.0	99	21	AAH86210	Human single nucle
C 145	10	100.0	51	22	AAH86210	Human single nucle	C 218	10	100.0	102	21	AAH86210	Human single nucle
C 146	10	100.0	51	22	AAH86210	Human single nucle	C 219	10	100.0	102	21	AAH86210	Human single nucle
C 147	10	100.0	51	22	AAH86210	Human single nucle	C 220	10	100.0	102	22	AAH86210	Human single nucle
C 148	10	100.0	51	23	ABL00766	DNA 53mer circle s	C 221	10	100.0	102	22	AAH86210	Human single nucle
C 149	10	100.0	53	19	AAH86210	Human single nucle	C 222	10	100.0	102	22	AAH86210	Human single nucle
C 150	10	100.0	53	19	AAH86210	Human single nucle	C 223	10	100.0	102	22	AAH86210	Human single nucle
C 151	10	100.0	53	19	AAH86210	Human single nucle	C 224	10	100.0	102	22	AAH86210	Human single nucle
C 152	10	100.0	53	19	AAH86210	Human single nucle	C 225	10	100.0	102	22	AAH86210	Human single nucle
C 153	10	100.0	53	20	AAH86210	Human single nucle	C 226	10	100.0	102	22	AAH86210	Human single nucle
C 154	10	100.0	54	17	AAH86210	Human single nucle	C 227	10	100.0	102	24	AAH86210	Human single nucle

C 228	10	100.0	103	21	AA42691	Human secreted exp	301	10	100.0	146	22	AAK19330	Human brain expres
C 229	10	100.0	104	21	AAC25734	Human secreted pro	302	10	100.0	146	22	AAK45307	Human bone marrow
C 230	10	100.0	105	22	ABA75693	Human foetal liver	303	10	100.0	146	22	AAI51261	Probe #19947 used
C 231	10	100.0	106	22	ABF83563	B. gymnorhiza sal	c 304	10	100.0	146	22	AAQ30393	Mouse brain p69 5'
C 232	10	100.0	107	24	ABL78374	Human ovarian canc	305	10	100.0	146	22	ABS44986	Human liver single
C 233	10	100.0	108	25	ABX55439	Bovine EST associa	306	10	100.0	146	24	ABS19584	Human genome-deriv
C 234	10	100.0	112	17	AAT39029	PCR primer 593 for	c 307	10	100.0	147	21	AAC23129	Human secreted pro
C 235	10	100.0	114	22	ABA73004	Human foetal liver	c 308	10	100.0	147	22	AAS34332	Human cDNA encodin
C 236	10	100.0	114	22	ABA74146	Human foetal liver	c 309	10	100.0	147	24	ABK85007	DNA encoding cadhe
C 237	10	100.0	114	22	ABA93160	Probe #17626 for g	c 310	10	100.0	147	24	ABK29624	Colon adenocarcino
C 238	10	100.0	114	22	AAK21436	Human brain expres	311	10	100.0	148	21	AAC19931	Human secreted pro
C 239	10	100.0	114	22	AAK22600	Human brain expres	312	10	100.0	148	21	AAC20151	Human secreted pro
C 240	10	100.0	114	22	AAK47597	Human bone marrow	314	10	100.0	149	22	AAI68590	S. tuberculosis SUT1
C 241	10	100.0	114	22	AAK48768	Human bone marrow	315	10	100.0	150	19	AAK10878	Human biallelic po
C 242	10	100.0	114	22	AAI53429	Probe #22115 used	c 316	10	100.0	150	19	AAK10879	Human biallelic po
C 243	10	100.0	114	22	AAI54598	Probe #23284 used	317	10	100.0	150	19	AAK12142	Human biallelic po
C 244	10	100.0	114	23	ABS47332	Human liver single	318	10	100.0	150	19	AAK10503	Human biallelic po
C 245	10	100.0	114	23	ABS48444	Human liver single	319	10	100.0	150	19	AAK10504	Human biallelic po
C 246	10	100.0	114	24	ABS21681	Human genome-deriv	c 320	10	100.0	152	21	AAC04835	Human secreted pro
C 247	10	100.0	114	24	ABS22480	Human genome-deriv	321	10	100.0	155	21	AAA43003	Human secreted exp
C 248	10	100.0	116	16	AAT22045	Human gene signatu	c 322	10	100.0	156	22	AAH84243	Human secreted pro
C 249	10	100.0	117	22	AAH84245	Human cell death p	c 323	10	100.0	156	24	ABK76880	Human cell death p
C 250	10	100.0	118	22	ABAI8306	Human nervous syst	324	10	100.0	156	24	ABN70111	Bacillus lichenifo
C 251	10	100.0	118	24	ABT03406	Ovary cell-specifi	c 325	10	100.0	157	22	AAI61613	Streptococcus poly
C 252	10	100.0	119	25	ABX30673	Human GDP-mannose	326	10	100.0	159	21	AAI61613	Streptococcus poly
C 253	10	100.0	121	22	ABA77351	P53 mutation corre	c 327	10	100.0	159	21	AAI61613	Soybean 318013 reg
C 254	10	100.0	121	22	ABA77352	P53 mutation corre	328	10	100.0	159	21	AAI61613	Fusarium venenatum
C 255	10	100.0	121	22	ABA77352	P53 mutation corre	329	10	100.0	161	22	AAK74123	Human immune/haema
C 256	10	100.0	121	22	ABA77368	P53 mutation corre	330	10	100.0	161	22	AAK74124	Human immune/haema
C 257	10	100.0	121	22	ABA77471	P53 mutation corre	331	10	100.0	161	22	AAK74125	Human immune/haema
C 258	10	100.0	121	22	ABA77472	P53 mutation corre	332	10	100.0	162	21	AAK66665	Novel human polynu
C 259	10	100.0	121	22	ABA77475	P53 mutation corre	c 333	10	100.0	163	25	ABZ78829	Human secreted pro
C 260	10	100.0	121	22	ABA77476	P53 mutation corre	334	10	100.0	163	24	ABV96596	Human pancreatic c
C 261	10	100.0	123	22	AAI03420	Human reproductive	c 335	10	100.0	163	25	ABZ78829	Human pancreatic c
C 262	10	100.0	123	22	AAI03421	Human reproductive	336	10	100.0	164	16	ABZ09376	Tumour suppression
C 263	10	100.0	124	21	AAA41960	Human secreted exp	337	10	100.0	164	16	AAI26273	Human oligonucleot
C 264	10	100.0	125	16	AAT05584	Mouse brain p69 cd	338	10	100.0	164	16	AAI21185	Human gene signatu
C 265	10	100.0	125	16	AAC09800	Human secreted pro	339	10	100.0	169	15	AAQ76860	Human gene signatu
C 266	10	100.0	125	24	ABS72812	Human gene trapped	340	10	100.0	169	25	AAQ76860	Human gene signatu
C 267	10	100.0	125	24	ABS72812	Human gene trapped	341	10	100.0	169	25	AAQ76860	Human gene signatu
C 268	10	100.0	126	20	AAI82264	Corn tassal-deriv	c 342	10	100.0	170	22	AAH83122	Human ovarian PCR-
C 269	10	100.0	126	20	AAI82264	Modified intron (I	343	10	100.0	170	22	AAH83122	Human ovarian PCR-
C 270	10	100.0	128	21	AAH84244	Human cell death p	c 344	10	100.0	172	21	AAK26682	Human gene trapped
C 271	10	100.0	128	21	AAH84244	Human cell death p	345	10	100.0	172	21	AAK26682	Human gene trapped
C 272	10	100.0	129	22	AAK24208	Human prostate can	c 346	10	100.0	174	22	AAI55976	Human secreted pro
C 273	10	100.0	129	22	AAI56202	Human brain expres	c 347	10	100.0	174	22	AAI55976	Human secreted pro
C 274	10	100.0	129	23	ABS49882	Probe #24888 used	c 348	10	100.0	176	22	AAK27878	Human cell death p
C 275	10	100.0	129	24	ABS49882	Human liver single	c 349	10	100.0	177	22	AAK27878	Human cell death p
C 276	10	100.0	132	20	AAI55702	Human genome-deriv	c 350	10	100.0	178	22	AAK72814	Human immune/haema
C 277	10	100.0	132	22	ABA70763	DNA sequence encod	c 351	10	100.0	178	23	ABK41929	Human immune/haema
C 278	10	100.0	132	22	ABA70763	Human foetal liver	c 352	10	100.0	180	24	ABN70012	cDNA encoding nove
C 279	10	100.0	132	22	AAK19016	Probe #15724 for g	353	10	100.0	181	22	AAI61656	Streptococcus poly
C 280	10	100.0	132	22	AAK44967	Human brain expres	354	10	100.0	182	24	ABU63623	Soybean 318013 reg
C 281	10	100.0	132	22	AAI56202	Probe #24888 used	355	10	100.0	182	24	ABU63623	Soybean 318013 reg
C 282	10	100.0	132	22	AAI56202	Probe #24888 used	356	10	100.0	183	22	ABA70927	Breast cancer rela
C 283	10	100.0	132	22	AAI56202	Probe #24888 used	357	10	100.0	183	22	ABA70927	Breast cancer rela
C 284	10	100.0	132	22	AAI56202	Probe #24888 used	358	10	100.0	183	22	AAI51115	Human foetal liver
C 285	10	100.0	132	22	AAI56202	Probe #24888 used	359	10	100.0	183	22	AAI51115	Human foetal liver
C 286	10	100.0	132	22	AAI56202	Probe #24888 used	360	10	100.0	183	22	AAI51115	Human foetal liver
C 287	10	100.0	132	22	AAI56202	Probe #24888 used	361	10	100.0	183	22	AAI51115	Human foetal liver
C 288	10	100.0	132	22	AAI56202	Probe #24888 used	362	10	100.0	183	22	AAI51115	Human foetal liver
C 289	10	100.0	132	22	AAI56202	Probe #24888 used	363	10	100.0	183	22	AAI51115	Human foetal liver
C 290	10	100.0	132	22	AAI56202	Probe #24888 used	364	10	100.0	183	22	AAI51115	Human foetal liver
C 291	10	100.0	132	22	AAI56202	Probe #24888 used	365	10	100.0	183	22	AAI51115	Human foetal liver
C 292	10	100.0	132	22	AAI56202	Probe #24888 used	366	10	100.0	183	22	AAI51115	Human foetal liver
C 293	10	100.0	132	22	AAI56202	Probe #24888 used	367	10	100.0	183	22	AAI51115	Human foetal liver
C 294	10	100.0	132	22	AAI56202	Probe #24888 used	368	10	100.0	183	22	AAI51115	Human foetal liver
C 295	10	100.0	132	22	AAI56202	Probe #24888 used	369	10	100.0	183	22	AAI51115	Human foetal liver
C 296	10	100.0	132	22	AAI56202	Probe #24888 used	370	10	100.0	183	22	AAI51115	Human foetal liver
C 297	10	100.0	132	22	AAI56202	Probe #24888 used	371	10	100.0	183	22	AAI51115	Human foetal liver
C 298	10	100.0	132	22	AAI56202	Probe #24888 used	372	10	100.0	183	22	AAI51115	Human foetal liver
C 299	10	100.0	132	22	AAI56202	Probe #24888 used	373	10	100.0	183	22	AAI51115	Human foetal liver
C 300	10	100.0	132	22	AAI56202	Probe #24888 used	374	10	100.0	183	22	AAI51115	Human foetal liver

C 374	10	100.0	10	100.0	10	100.0	218	22	AAI21506	Probe #11439 for g
C 375	10	100.0	10	100.0	10	100.0	218	22	AAI46798	Probe #15484 used
C 376	10	100.0	10	100.0	10	100.0	218	22	AAI07202	Probe #7193 used t
C 377	10	100.0	10	100.0	10	100.0	218	22	ABS40316	Human liver single
C 378	10	100.0	10	100.0	10	100.0	218	23	ABS14697	Human genome-deriv
C 379	10	100.0	10	100.0	10	100.0	218	24	ABL87111	Human ovarian carc
C 380	10	100.0	10	100.0	10	100.0	218	24	AAI55455	Human genome-deriv
C 381	10	100.0	10	100.0	10	100.0	218	24	ABS49139	Human secreted exp
C 382	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 383	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 384	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 385	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
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C 388	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 389	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 390	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 391	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 392	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
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C 399	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 400	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 401	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 402	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 403	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 404	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 405	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 406	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 407	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
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C 427	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 428	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 429	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
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C 435	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
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C 439	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 440	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 441	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 442	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 443	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 444	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 445	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp
C 446	10	100.0	10	100.0	10	100.0	218	24	ABS23061	Human secreted exp

ALIGNMENTS

RESULT 1

AAV50115

ID AAV50115 standard; DNA; 10 BP.

XX

AC AAV50115;

XX

DT 21-OCT-1998 (first entry)

XX

XX Yeast tag for NORF gene locus NORF5.

XX

XX Yeast: Saccharomyces cerevisiae; transcriptome; cell cycle;

XX regulation; eukaryotic cell; antifungal; SAGE tag; gene expression;

XX serial analysis of gene expression; probe; ss.

XX

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OS Saccharomyces cerevisiae.
OS Synthetic.
XX
XX
PN WO9832847-A2.
XX
XX
PD 30-JUL-1998.
XX
XX
PF 22-JAN-1998; 98WO-US01216.
XX
XX
PR 23-JAN-1997; 97US-0035917.
XX
XX
PA (UYJO ) UNIV JOHNS HOPKINS SCHOOL MEDICINE.
XX
XX
PI Kinzler KW, Velculescu VE, Vogelstein B;
XX
XX
DR WPI; 1998-427943/36.
XX
XX
PT Yeast transcriptome - useful for modulating eukaryotic cell, for
PT screening antifungal agents, and for identifying genes in cell cycle
PT progression
XX
XX
PS Claim 1; Page 23; 44pp; English.
XX
XX
CC Yeast transcriptome is encoded by a DNA molecule comprising a yeast
CC gene involved in cell cycle progression selected from the group of
CC nonannotated ORF (NORF) genes. SAGE (serial analysis of gene expression)
CC tags for highly expressed genes and NORF genes are given in AAV50051 to
CC AAV50345. The present invention describes: (1) a method of using yeast
CC genes to modulate the cell cycle which comprises administering to a cell
CC an isolated DNA molecule comprising a yeast gene which is involved in
CC cell cycle progression selected from differentially expressed genes
CC (SAGE tags given in AAV50051 to AAV50345); (2) a method for screening
CC candidate antifungal drugs which comprises contacting a test substance
CC with a yeast cell and monitoring expression of a yeast gene which is
CC involved in cell cycle progression; (3) a method of identifying human
CC genes which are involved in cell cycle progression which comprises
CC hybridizing a probe comprising at least 10 contiguous nucleotides of a
CC yeast gene which is differentially expressed between at least 2 phases
CC selected from the log phase, the S phase and the G2/M phase; and (4) a
CC probe for ascertaining the phase in the cell cycle, where the probe
CC comprises at least 14 contiguous nucleotides of a NORF gene (SAGE tags
CC given in AAV50051 to AAV50345), or as an array of probes on a solid
CC support.
XX
SQ Sequence 10 BP; 0 A; 3 C; 0 G; 7 T; 0 other;
Query Match 100.0%; Score 10; DB 19; Length 10;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 1 CTTCTCTTTT 10
RESULT 2
AAF33332
ID AAF33332 standard; DNA; 10 BP.
XX
XX
AC AAF33332;
XX
XX
DT 23-MAR-2001 (first entry)
XX
DE Yeast NORF gene SAGE tag oligonucleotide SEQ ID NO:71.
XX
XX
KW Yeast; Saccharomyces cerevisiae; characterisation; cell cycle; NORF;
KW nor previously assigned open reading frame; nonannotated ORF; SAGE;
KW serial analysis of gene expression; antifungal; tag; identification;
KW linker; PCR primer; ds.
XX
OS Saccharomyces cerevisiae.
XX
XX
PN WO200077214-A2.
XX
XX

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XX
XX
PD 21-DEC-2000.
XX
XX
PF 14-JUN-2000; 2000WO-US16223.
XX
XX
PR 16-JUN-1999; 99US-0335032.
XX
XX
PA (UYJO ) UNIV JOHNS HOPKINS.
XX
XX
PI Velculescu V, Vogelstein B, Kinzler K;
XX
XX
DR WPI; 2001-061874/07.
XX
XX
PT Yeast gene coding sequences comprising NORF genes with serial analysis
PT of gene expression (SAGE) tags, useful for studying, monitoring and
PT affecting phases of the cell cycle.
XX
XX
PS Claim 1; Page 23; 419pp; English.
XX
XX
CC The present invention describes an isolated DNA molecule comprising a
CC coding sequence of a yeast gene selected from a group of 745 NORF (not
CC previously assigned open reading frame; or nonannotated ORF) genes
CC comprising a SAGE (serial analysis of gene expression) tag. Also
CC described are: (1) a method (M1) of using NORF genes to affect the cell
CC cycle comprising administering a NORF gene whose expression varies by at
CC least 10% between any two phases of the cell cycle selected from log
CC phase, S phase and G2/M; (2) a method (M2) for screening candidate
CC antifungal drugs comprising: (a) contacting a test substance with a
CC yeast cell; and (b) monitoring expression of a NORF gene whose
CC expression varies as in M1, where a test substance which modifies the
CC expression of the yeast gene is a candidate antifungal drug; (3) a method
CC (M3) for identifying human genes which are involved in cell cycle
CC progression comprising contacting human DNA with a probe which comprises
CC at least 10 contiguous nucleotides of a NORF gene whose expression varies
CC as in M1; and (4) a method (M4) for identifying a candidate drug as a
CC member of a class of drugs having a characteristic effect on gene
CC expression in a yeast cell comprising contacting a yeast cell with a
CC candidate drug and monitoring expression in the yeast cell of at least 1
CC NORF gene whose expression is affected by the class of drugs. The NORF
CC genes may be used to study, monitor and affect phases of the cell cycle,
CC the differentially expressed genes may be used as markers of phases of
CC the cell cycle. The methods may be used to identify candidate drugs which
CC affect the cell cycle and for identification of antifungal drugs.
CC AAF33268 to AAF4064 represent SAGE tags used in the exemplification of
CC the present invention. AAF33262 to AAF33267 represent linkers and PCR
CC primers used in the SAGE method, in the exemplification of the present
CC invention.
XX
SQ Sequence 10 BP; 0 A; 3 C; 0 G; 7 T; 0 other;
Query Match 100.0%; Score 10; DB 22; Length 10;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 1 CTTCTCTTTT 10
RESULT 3
AAF33321
ID AAF33321 standard; DNA; 10 BP.
XX
XX
AC AAF33321;
XX
XX
DT 23-MAR-2001 (first entry)
XX
DE Yeast NORF gene SAGE tag oligonucleotide SEQ ID NO:660.
XX
XX
KW Yeast; Saccharomyces cerevisiae; characterisation; cell cycle; NORF;
KW nor previously assigned open reading frame; nonannotated ORF; SAGE;
KW serial analysis of gene expression; antifungal; tag; identification;
KW linker; PCR primer; ds.
XX

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XX OS Saccharomyces cerevisiae.
 XX PN WO200077214-A2.
 XX PD 21-DEC-2000.
 XX XX 14-JUN-2000; 2000WO-US16223.
 XX PF 16-JUN-1999; 99US-0335032.
 XX PR (UYJO) UNIV JOHNS HOPKINS.
 XX PA Velculescu V, Vogelstein B, Kinzler K;
 XX DR WPI; 2001-061874/07.
 XX XX Yeast gene coding sequences comprising NORF genes with serial analysis
 PT of gene expression (SAGE) tags, useful for studying, monitoring and
 PT affecting phases of the cell cycle -
 XX
 PS Claim 1; Page 398; 419pp; English.
 XX
 CC The present invention describes an isolated DNA molecule comprising a
 CC coding sequence of a yeast gene selected from a group of 745 NORF (not
 CC previously assigned open reading frame; or nonannotated ORF) genes
 CC comprising a SAGE (serial analysis of gene expression) tag. Also
 CC described are: (1) a method (M1) of using NORF genes to affect the cell
 CC cycle comprising administering a NORF gene whose expression varies by at
 CC least 10% between any two phases of the cell cycle selected from log
 CC phase, S phase and G2/M; (2) a method (M2) for screening candidate
 CC antifungal drugs comprising: (a) contacting a test substance with a
 CC yeast cell; and (b) monitoring expression of a NORF gene whose
 CC expression varies as in M1, where a test substance which modifies the
 CC expression of the yeast gene is a candidate antifungal drug; (3) a method
 CC (M3) for identifying human genes which are involved in cell cycle
 CC progression comprising contacting human DNA with a probe which comprises
 CC at least 10 contiguous nucleotides of a NORF gene whose expression varies
 CC as in M1; and (4) a method (M4) for identifying a candidate drug as a
 CC member of a class of drugs having a characteristic effect on gene
 CC expression in a yeast cell comprising contacting a yeast cell with a
 CC candidate drug and monitoring expression in the yeast cell of at least 1
 CC NORF gene whose expression is affected by the class of drugs. The NORF
 CC genes may be used to study, monitor and affect phases of the cell cycle,
 CC the differentially expressed genes may be used as markers of phases of
 CC the cell cycle. The methods may be used to identify candidate drugs which
 CC affect the cell cycle and for identification of antifungal drugs.
 CC AAF33268 to AAF44064 represent SAGE tags used in the exemplification of
 CC the present invention. AAF33262 to AAF33267 represent linkers and PCR
 CC primers used in the SAGE method, in the exemplification of the present
 CC invention.
 XX
 SQ Sequence 10 BP; 0 A; 3 C; 0 G; 7 T; 0 other;
 Query Match 100.0%; Score 10; DB 22; Length 10;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 1 CTTCTCTTTT 10
 RESULT 4
 AAF33924
 ID AAF33924 standard; DNA; 10 BP.
 XX
 XX AAF33924;
 AC
 DT 23-MAR-2001 (first entry)
 XX
 DE Yeast NORF gene SAGE tag oligonucleotide SEQ ID NO:663.
 XX

KW Yeast; Saccharomyces cerevisiae; characterisation; cell cycle; NORF;
 KW nor previously assigned open reading frame; nonannotated ORF; SAGE;
 KW serial analysis of gene expression; antifungal; tag; identification;
 KW linker; PCR primer; ds.
 XX
 OS Saccharomyces cerevisiae.
 XX PN WO200077214-A2.
 XX PD 21-DEC-2000.
 XX XX 14-JUN-2000; 2000WO-US16223.
 XX PF 16-JUN-1999; 99US-0335032.
 XX PR (UYJO) UNIV JOHNS HOPKINS.
 XX PA Velculescu V, Vogelstein B, Kinzler K;
 XX DR WPI; 2001-061874/07.
 XX XX Yeast gene coding sequences comprising NORF genes with serial analysis
 PT of gene expression (SAGE) tags, useful for studying, monitoring and
 PT affecting phases of the cell cycle -
 XX
 PS Claim 1; Page 398; 419pp; English.
 XX
 CC The present invention describes an isolated DNA molecule comprising a
 CC coding sequence of a yeast gene selected from a group of 745 NORF (not
 CC previously assigned open reading frame; or nonannotated ORF) genes
 CC comprising a SAGE (serial analysis of gene expression) tag. Also
 CC described are: (1) a method (M1) of using NORF genes to affect the cell
 CC cycle comprising administering a NORF gene whose expression varies by at
 CC least 10% between any two phases of the cell cycle selected from log
 CC phase, S phase and G2/M; (2) a method (M2) for screening candidate
 CC antifungal drugs comprising: (a) contacting a test substance with a
 CC yeast cell; and (b) monitoring expression of a NORF gene whose
 CC expression varies as in M1, where a test substance which modifies the
 CC expression of the yeast gene is a candidate antifungal drug; (3) a method
 CC (M3) for identifying human genes which are involved in cell cycle
 CC progression comprising contacting human DNA with a probe which comprises
 CC at least 10 contiguous nucleotides of a NORF gene whose expression varies
 CC as in M1; and (4) a method (M4) for identifying a candidate drug as a
 CC member of a class of drugs having a characteristic effect on gene
 CC expression in a yeast cell comprising contacting a yeast cell with a
 CC candidate drug and monitoring expression in the yeast cell of at least 1
 CC NORF gene whose expression is affected by the class of drugs. The NORF
 CC genes may be used to study, monitor and affect phases of the cell cycle,
 CC the differentially expressed genes may be used as markers of phases of
 CC the cell cycle. The methods may be used to identify candidate drugs which
 CC affect the cell cycle and for identification of antifungal drugs.
 CC AAF33268 to AAF44064 represent SAGE tags used in the exemplification of
 CC the present invention. AAF33262 to AAF33267 represent linkers and PCR
 CC primers used in the SAGE method, in the exemplification of the present
 CC invention.
 XX
 SQ Sequence 10 BP; 0 A; 3 C; 0 G; 7 T; 0 other;
 Query Match 100.0%; Score 10; DB 22; Length 10;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 1 CTTCTCTTTT 10
 RESULT 5
 AAF34233
 ID AAF34233 standard; DNA; 10 BP.
 XX
 XX AAF34233;
 AC
 XX


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OS Micrococcus luteus.
PN US5861244-A.
XX
XX
PD 19-JAN-1999.
XX
XX 22-DEC-1993; 93US-0173489.
XX
XX 22-DEC-1993; 93US-0173489.
PR 29-OCT-1992; 92US-0969436.
XX
XX (PROF-) PROFILE DIAGNOSTIC SCI INC.
XX
XX Hepburn AG, Wang C;
XX
XX WPI; 1999-130384/11.
XX
XX Assay of genetic sequences based on triplex formation from double
PT stranded analyte - and hybrid of anchor and reporter sequences, with
PT reporter released if triplex formation occurs, used e.g. to identify
PT bacteria
XX
XX Disclosure; Columns 21-22; 168pp; English.
XX
XX The present sequence represents a polynucleotide that is able to
CC form a triple helix with a double stranded sequence. Cytosine bases
CC in the present can be replaced with 5-methylcytosine for increased
CC triplex stability. The present sequence is used in the assay of the
CC invention, where it can be part of the anchor DNA or reporter DNA
CC sequence. The assay comprises adding a sample containing double-stranded
CC DNA test sequences to an aqueous medium containing at least one complex
CC of anchor DNA, attached to a solid support, and reporter DNA, where
CC either a part of the anchor DNA or reporter DNA is designed to form
CC a triple-strand structure with part of the test sequence. Triplex
CC formation results in displacement of the reporter DNA which is
CC detected as an indication of the presence of the DNA test sequence.
CC The method is used to detect DNA sequences, particularly for
CC identification of bacteria (by detecting genes for ribosomal RNA) in
CC clinical samples, but also detection of oncogenes and Hepatitis B virus.
XX
XX Sequence 12 BP; 0 A; 4 C; 0 G; 8 T; 0 other;

Query Match 100.0%; Score 10; DB 20; Length 12;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 3 CTTCTCTTTT 12

RESULT 8
ABH67817
ID ABH67817 standard; DNA; 12 BP.
XX
XX ABH67817;
AC
XX 22-FEB-2002 (first entry)
XX
XX Oligonucleotide primer SEQ ID NO 267794 for detecting SNP TSC0000531.
DE
DE SNP: single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
XX Homo sapiens.
OS
XX WO200177384-A2.
PN
XX 18-OCT-2001.
PD
XX 06-APR-2001; 2001WO-IB00713.
PF

PR 07-APR-2000; 2000DE-1019173.
XX
XX (EPIG-) EPIGENOMICS AG.
PI
XX Olek A, Piepenbrock C, Berlin K;
XX
XX WPI; 2001-657177/75.
DR
XX Set of oligonucleotides, useful for diagnosis and cell typing, is
PT designed to detect single nucleotide polymorphisms and cytosine
PT methylation status -
XX
XX Claim 1; SEQ ID 267794; 29pp + Sequence Listing; German.
XX
XX This invention describes novel oligonucleotide primers or peptide nucleic
CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
CC and cytosine methylation status in chemically pretreated genomic DNA. The
CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
CC range of diseases including immune system, gastrointestinal, respiratory,
CC central nervous system, cardiovascular and metabolic disorders. The
CC oligomers are also used for detecting cell type differentiation.
CC ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and
CC ABT00010-ABT99989 represent the oligomers described in the invention.
CC NOTE: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 12 BP; 1 A; 4 C; 0 G; 7 T; 0 other;

Query Match 100.0%; Score 10; DB 23; Length 12;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 2 CTTCTCTTTT 11

RESULT 9
ABI80052
ID ABI80052 standard; DNA; 12 BP.
XX
XX ABI80052;
AC
XX 22-FEB-2002 (first entry)
XX
XX Oligonucleotide primer SEQ ID NO 380025 for detecting SNP TSC0063600.
DE
DE SNP: single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
XX Homo sapiens.
OS
XX WO200177384-A2.
PN
XX 18-OCT-2001.
PD
XX 06-APR-2001; 2001WO-IB00713.
PF
XX 07-APR-2000; 2000DE-1019173.
XX
XX (EPIG-) EPIGENOMICS AG.
PA
XX Olek A, Piepenbrock C, Berlin K;
PI
XX WPI; 2001-657177/75.
DR
XX Set of oligonucleotides, useful for diagnosis and cell typing, is
PT designed to detect single nucleotide polymorphisms and cytosine
PT methylation status -
XX
XX Claim 1; SEQ ID 380025; 29pp + Sequence Listing; German.
PS

```


XX This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
 CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation.
 CC ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and
 CC AB100010-AB182073 represent the oligomers described in the invention.
 CC NOTE: The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.

XX SQ Sequence 12 BP; 1 A; 4 C; 0 G; 7 T; 0 other;

Query Match 100.0%; Score 10; DB 23; Length 12;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
 Db 3 CTTCTCTTTT 12

RESULT 10
 ABC07094/c
 ID ABC07094 standard; DNA; 13 BP.
 XX AC ABC07094;
 XX DT 20-FEB-2002 (first entry)
 XX DE Oligonucleotide SEQ ID NO 7085 for detecting SNP TSC00002097.
 XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
 KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
 KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
 XX OS Homo sapiens.
 XX PN WO200177384-A2.
 XX PD 18-OCT-2001.
 XX PF 06-APR-2001; 2001WO-IB00713.
 XX PR 07-APR-2000; 2000DE-1019173.
 XX PA (EPIG-) EPIGENOMICS AG.
 XX PI Olek A, Piepenbrock C, Berlin K;
 XX WPI; 2001-657177/75.
 XX Set of oligonucleotides, useful for diagnosis and cell typing, is
 PT designed to detect single nucleotide polymorphisms and cytosine
 PT methylation status -
 XX Claim 1; SEQ ID 7085; 29pp + Sequence Listing; German.

This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
 CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation.
 CC ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and
 CC AB100010-AB182073 represent the oligomers described in the invention.
 CC NOTE: The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.

XX SQ Sequence 13 BP; 8 A; 0 C; 4 G; 1 T; 0 other;

Query Match 100.0%; Score 10; DB 23; Length 13;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
 Db 13 CTTCTCTTTT 4

RESULT 11
 ABC07095
 ID ABC07095 standard; DNA; 13 BP.
 XX AC ABC07095;
 XX DT 20-FEB-2002 (first entry)
 XX DE Oligonucleotide SEQ ID NO 7086 for detecting SNP TSC00002097.
 XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
 KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
 KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
 XX OS Homo sapiens.
 XX PN WO200177384-A2.
 XX PD 18-OCT-2001.
 XX PF 06-APR-2001; 2001WO-IB00713.
 XX PR 07-APR-2000; 2000DE-1019173.
 XX PA (EPIG-) EPIGENOMICS AG.
 XX PI Olek A, Piepenbrock C, Berlin K;
 XX WPI; 2001-657177/75.
 XX Set of oligonucleotides, useful for diagnosis and cell typing, is
 PT designed to detect single nucleotide polymorphisms and cytosine
 PT methylation status -
 XX Claim 1; SEQ ID 7086; 29pp + Sequence Listing; German.

This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
 CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation.
 CC ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and
 CC AB100010-AB182073 represent the oligomers described in the invention.
 CC NOTE: The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.

XX SQ Sequence 13 BP; 1 A; 4 C; 0 G; 8 T; 0 other;

Query Match 100.0%; Score 10; DB 23; Length 13;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
 Db 1 CTTCTCTTTT 10

RESULT 12

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ABC53772/c
ID ABC53772 standard; DNA; 13 BP.
XX
AC ABC53772;
XX
XX 21-FEB-2002 (first entry)
XX
XX Oligonucleotide SEQ ID NO 53789 for detecting SNP TSC0014813.
XX
XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
XX peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
XX central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
XX Homo sapiens.
XX
XX WO200177384-A2.
XX
XX 18-OCT-2001.
XX
XX 06-APR-2001; 2001WO-IB00713.
XX
XX 07-APR-2000; 2000DE-1019173.
XX
XX (EPIG-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K;
XX
XX WPI; 2001-657177/75.
XX
XX Set of oligonucleotides, useful for diagnosis and cell typing, is
XX designed to detect single nucleotide polymorphisms and cytosine
XX methylation status -
XX
XX Claim 1; SEQ ID 53789; 29pp + Sequence Listing; German.
XX
XX This invention describes novel oligonucleotide primers or peptide nucleic
XX acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
XX and cytosine methylation status in chemically pretreated genomic DNA. The
XX oligonucleotides are used for diagnosis and/or prognosis of cancer and a
XX range of diseases including immune system, gastrointestinal, respiratory,
XX central nervous system, cardiovascular and metabolic disorders. The
XX oligomers are also used for detecting cell type differentiation.
XX ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and
XX ABI00010-ABI82073 represent the oligomers described in the invention.
XX NOTE: The sequence data for this patent did not form part of the printed
XX specification, but was obtained in electronic format from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 13 BP; 10 A; 0 C; 3 G; 0 U; 0 other;
XX
XX Query Match 100.0%; Score 10; DB 23; Length 13;
XX Best Local Similarity 100.0%; Pred. No. 1.8e+04;
XX Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 CTTCTCTTTT 10
XX Db 10 CTTCTCTTTT 1
XX
XX RESULT 13
XX ABC53773
XX ID ABC53773 standard; DNA; 13 BP.
XX
XX AC ABC53773;
XX
XX 21-FEB-2002 (first entry)
XX
XX Oligonucleotide SEQ ID NO 53790 for detecting SNP TSC0014813.
XX
XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
XX peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
XX central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
XX Homo sapiens.
XX
XX WO200177384-A2.
XX
XX 18-OCT-2001.
XX
XX 06-APR-2001; 2001WO-IB00713.
XX
XX 07-APR-2000; 2000DE-1019173.
XX
XX (EPIG-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K;
XX
XX WPI; 2001-657177/75.
XX
XX Set of oligonucleotides, useful for diagnosis and cell typing, is
XX designed to detect single nucleotide polymorphisms and cytosine
XX methylation status -
XX
XX Claim 1; SEQ ID 53789; 29pp + Sequence Listing; German.
XX
XX This invention describes novel oligonucleotide primers or peptide nucleic
XX acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP)
XX and cytosine methylation status in chemically pretreated genomic DNA. The
XX oligonucleotides are used for diagnosis and/or prognosis of cancer and a
XX range of diseases including immune system, gastrointestinal, respiratory,
XX central nervous system, cardiovascular and metabolic disorders. The
XX oligomers are also used for detecting cell type differentiation.
XX ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and
XX ABI00010-ABI82073 represent the oligomers described in the invention.
XX NOTE: The sequence data for this patent did not form part of the printed
XX specification, but was obtained in electronic format from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 13 BP; 10 A; 0 C; 3 G; 0 U; 0 other;
XX
XX Query Match 100.0%; Score 10; DB 23; Length 13;
XX Best Local Similarity 100.0%; Pred. No. 1.8e+04;
XX Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 CTTCTCTTTT 10
XX Db 10 CTTCTCTTTT 1
XX
XX RESULT 14
XX ABH25906/c
XX ID ABH25906 standard; DNA; 13 BP.
XX
XX AC ABH25906;
XX
XX 22-FEB-2002 (first entry)
XX
XX Oligonucleotide SEQ ID NO 225883 for detecting SNP TSC0055065.
XX
XX SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
XX peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
XX central nervous system; gastrointestinal; respiratory; immune; metabolic.
XX
XX Homo sapiens.
XX
XX WO200177384-A2.
XX
XX 18-OCT-2001.
XX
XX 06-APR-2001; 2001WO-IB00713.
XX
XX 07-APR-2000; 2000DE-1019173.
XX
XX (EPIG-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K;

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XX WPI; 2001-657177/75.
 DR Set of oligonucleotides, useful for diagnosis and cell typing, is
 PT designed to detect single nucleotide polymorphisms and cytosine
 PT methylation status -
 PS Claim 1; SEQ ID 225883; 29pp + Sequence Listing; German.
 XX This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP).
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a
 CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation.
 CC ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and
 CC ABI00010-ABI82073 represent the oligomers described in the invention.
 CC NOTE: The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.
 XX Sequence 13 BP; 8 A; 0 C; 3 G; 2 T; 0 other;
 SQ Query Match 100.0%; Score 10; DB 23; Length 13;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 10 CTTCTCTTTT 1
 RESULT 15
 ID ABH25907 standard; DNA; 13 BP.
 XX AC ABH25907;
 XX DT 22-FEB-2002 (first entry)
 XX OLigonucleotide SEQ ID NO 225884 for detecting SNP TSC0055065.
 DE SNP; single nucleotide polymorphism; human; diagnosis; PNA; cancer; CNS;
 KW peptide nucleic acid; cytosine methylation; cardiovascular; primer; ss;
 KW central nervous system; gastrointestinal; respiratory; immune; metabolic.
 XX Homo sapiens.
 OS WO200177384-A2.
 XX WO200177384-A2.
 XX 18-OCT-2001.
 XX 06-APR-2001; 2001WO-IB00713.
 XX 07-APR-2000; 2000DE-1019173.
 XX (EPIG-) EPIGENOMICS AG.
 XX Olek A, Piepenbrock C, Berlin K;
 XX WPI; 2001-657177/75.
 DR Set of oligonucleotides, useful for diagnosis and cell typing, is
 PT designed to detect single nucleotide polymorphisms and cytosine
 PT methylation status -
 PS Claim 1; SEQ ID 225884; 29pp + Sequence Listing; German.
 XX This invention describes novel oligonucleotide primers or peptide nucleic
 CC acid (PNA) oligomers for detecting single nucleotide polymorphisms (SNP).
 CC and cytosine methylation status in chemically pretreated genomic DNA. The
 CC oligonucleotides are used for diagnosis and/or prognosis of cancer and a

CC range of diseases including immune system, gastrointestinal, respiratory,
 CC central nervous system, cardiovascular and metabolic disorders. The
 CC oligomers are also used for detecting cell type differentiation.
 CC ABC00010-ABC99989, ABF00010-ABF99989, ABH00010-ABH99989 and
 CC ABI00010-ABI82073 represent the oligomers described in the invention.
 CC NOTE: The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.
 XX Sequence 13 BP; 2 A; 3 C; 0 G; 8 T; 0 other;
 SQ Query Match 100.0%; Score 10; DB 23; Length 13;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 4 CTTCTCTTTT 13
 RESULT 16
 ID ABV80474/c
 XX ID ABV80474 standard; DNA; 17 BP.
 XX AC ABV80474;
 XX DT 03-JAN-2003 (first entry)
 XX Human HTPL scanning oligonucleotide SEQ ID 1720.
 DE Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
 XX Homo sapiens.
 OS EPI229046-A2.
 XX EPI229046-A2.
 XX 07-AUG-2002.
 XX 28-JAN-2002; 2002EP-0001167.
 XX 30-JAN-2001; 2001WO-US000663.
 XX 30-JAN-2001; 2001WO-US000664.
 XX 30-JAN-2001; 2001WO-US000665.
 XX 30-JAN-2001; 2001WO-US000667.
 XX 30-JAN-2001; 2001WO-US000668.
 XX 30-JAN-2001; 2001WO-US000669.
 XX 23-MAY-2001; 2001US-0864761.
 XX 09-OCT-2001; 2001US-0327898.
 XX (AEON-) AEOMICA INC.
 XX Zhan J;
 XX WPI; 2002-676582/73.
 XX Novel isolated human testis expressed Patched like protein (HTPL),
 PT useful for identifying agonist and antagonist and specific binding
 PT partners, and for treating subjects having defects in HTPL -
 XX Example 2; Page 289; 718pp; English.
 XX The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and AB898519 to AB898520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
 CC shares an overall structure organisation with the Patched protein. The
 CC shared structural features strongly imply that HTPL plays a role similar
 CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was

CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC therapy and manufacture of a medicament for treatment or prevention of
 CC such disorder associated with decreased expression or activity of human
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.

XX SQ Sequence 17 BP; 8 A; 0 C; 7 G; 2 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||
 Db 17 CTTCTCTTTT 8

RESULT 17

ABV80475/c
 ID ABV80475 standard; DNA; 17 BP.

XX AC ABV80475;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 1721.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EPI229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US00663.

XX PR 30-JAN-2001; 2001WO-US00664.

XX PR 30-JAN-2001; 2001WO-US00665.

XX PR 30-JAN-2001; 2001WO-US00667.

XX PR 30-JAN-2001; 2001WO-US00668.

XX PR 30-JAN-2001; 2001WO-US00669.

XX PR 23-MAY-2001; 2001US-0864761.

XX PR 09-OCT-2001; 2001US-0327898.

XX PA (AEOM-) AEOMICA INC.

XX PI Zhan J;

XX DR WPI; 2002-676582/73.

XX PT Novel isolated human testis expressed Patched like protein (HTPL),
 PT useful for identifying agonist and antagonist and specific binding
 PT partners, and for treating subjects having defects in HTPL -

XX PS Example 2; Page 289; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
 CC shares an overall structure organisation with the Patched protein. The
 CC shared structural features strongly imply that HTPL plays a role similar

CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC therapy and manufacture of a medicament for treatment or prevention of
 CC such disorder associated with decreased expression or activity of human
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate, are
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.

XX SQ Sequence 17 BP; 8 A; 0 C; 6 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||
 Db 16 CTTCTCTTTT 7

RESULT 18

ABV80476/c

ID ABV80476 standard; DNA; 17 BP.

XX AC ABV80476;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 1722.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EPI229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US00663.

XX PR 30-JAN-2001; 2001WO-US00664.

XX PR 30-JAN-2001; 2001WO-US00665.

XX PR 30-JAN-2001; 2001WO-US00667.

XX PR 30-JAN-2001; 2001WO-US00668.

XX PR 23-MAY-2001; 2001US-0864761.

XX PR 09-OCT-2001; 2001US-0327898.

XX PA (AEOM-) AEOMICA INC.

XX PI Zhan J;

XX DR WPI; 2002-676582/73.

XX PT Novel isolated human testis expressed Patched like protein (HTPL),
 PT useful for identifying agonist and antagonist and specific binding
 PT partners, and for treating subjects having defects in HTPL -

XX PS Example 2; Page 289; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL

CC shares an overall structure organisation with the Patched protein. The
 CC shared structural features strongly imply that HTPL plays a role similar
 CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC therapy and manufacture of a medicament for treatment or prevention of
 CC such disorder associated with decreased expression or activity of human
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.

XX Sequence 17 BP; 9 A; 0 C; 5 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 Db 15 CTTCTCTTTT 6

RESULT 19

ABV80477/c
 ID ABV80477 standard; DNA; 17 BP.

XX AC ABV80477;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 1723.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EP1229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US00663.

XX PR 30-JAN-2001; 2001WO-US00664.

XX PR 30-JAN-2001; 2001WO-US00665.

XX PR 30-JAN-2001; 2001WO-US00667.

XX PR 30-JAN-2001; 2001WO-US00668.

XX PR 23-MAY-2001; 2001US-0864761.

XX PR 09-OCT-2001; 2001US-0327898.

XX PA (AEOM-) AEOMICA INC.

XX PI Zhan J;

XX DR WPI; 2002-676582/73.

XX PS Example 2; Page 289; 718pp; English.

XX The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the

CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
 CC shares an overall structure organisation with the Patched protein. The
 CC shared structural features strongly imply that HTPL plays a role similar
 CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC therapy and manufacture of a medicament for treatment or prevention of
 CC such disorder associated with decreased expression or activity of human
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.

XX Sequence 17 BP; 8 A; 0 C; 6 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 Db 14 CTTCTCTTTT 5

RESULT 20

ABV80478/c
 ID ABV80478 standard; DNA; 17 BP.

XX AC ABV80478;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 1724.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EP1229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US00663.

XX PR 30-JAN-2001; 2001WO-US00664.

XX PR 30-JAN-2001; 2001WO-US00665.

XX PR 30-JAN-2001; 2001WO-US00667.

XX PR 30-JAN-2001; 2001WO-US00668.

XX PR 23-MAY-2001; 2001US-0864761.

XX PR 09-OCT-2001; 2001US-0327898.

XX PA (AEOM-) AEOMICA INC.

XX PI Zhan J;

XX DR WPI; 2002-676582/73.

XX PS Novel isolated human testis expressed Patched like protein (HTPL),
 useful for identifying agonist and antagonist and specific binding
 partners, and for treating subjects having defects in HTPL -
 Example 2; Page 289; 718pp; English.

XX The present invention relates to human testis expressed Patched like

CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
 CC shares an overall structure organisation with the Patched protein. The
 CC shared structural features strongly imply that HTPL plays a role similar
 CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC therapy and manufacture of a medicament for treatment or prevention of
 CC such disorder associated with decreased expression or activity of human
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.

XX
 SQ Sequence 17 BP; 8 A; 0 C; 6 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||
 Db 13 CTTCTCTTTT 4

RESULT 21
 ABV80479/c
 ID ABV80479 standard; DNA; 17 BP.
 XX
 AC ABV80479;
 XX
 DT 03-JAN-2003 (first entry)
 XX
 DE Human HTPL scanning oligonucleotide SEQ ID 1725.
 XX
 KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
 XX
 OS Homo sapiens.
 XX
 PN EP1229046-A2.
 XX
 PD 07-AUG-2002.
 XX
 PF 28-JAN-2002; 2002EP-0001167.
 XX
 PR 30-JAN-2001; 2001WO-US00663.
 PR 30-JAN-2001; 2001WO-US00664.
 PR 30-JAN-2001; 2001WO-US00665.
 PR 30-JAN-2001; 2001WO-US00667.
 PR 30-JAN-2001; 2001WO-US00668.
 PR 30-JAN-2001; 2001WO-US00669.
 PR 23-MAY-2001; 2001US-0864761.
 PR 09-OCT-2001; 2001US-0327898.
 XX
 PA (AEOM-) AEOMICA INC.
 XX
 PI Zhan J;
 XX
 DR WPI; 2002-676582/73.
 XX
 PT Novel isolated human testis expressed Patched like protein (HTPL),
 PT useful for identifying agonist and antagonist and specific binding
 PT partners, and for treating subjects having defects in HTPL -
 XX
 PS Example 2; Page 290; 718pp; English.

XX
 CC The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
 CC shares an overall structure organisation with the Patched protein. The
 CC shared structural features strongly imply that HTPL plays a role similar
 CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC therapy and manufacture of a medicament for treatment or prevention of
 CC such disorder associated with decreased expression or activity of human
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.

XX
 SQ Sequence 17 BP; 8 A; 1 C; 5 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||
 Db 12 CTTCTCTTTT 3

RESULT 22
 ABV80480/c
 ID ABV80480 standard; DNA; 17 BP.
 XX
 AC ABV80480;
 XX
 DT 03-JAN-2003 (first entry)
 XX
 DE Human HTPL scanning oligonucleotide SEQ ID 1726.
 XX
 KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
 XX
 OS Homo sapiens.
 XX
 PN EP1229046-A2.
 XX
 PD 07-AUG-2002.
 XX
 PF 28-JAN-2002; 2002EP-0001167.
 XX
 PR 30-JAN-2001; 2001WO-US00663.
 PR 30-JAN-2001; 2001WO-US00664.
 PR 30-JAN-2001; 2001WO-US00665.
 PR 30-JAN-2001; 2001WO-US00667.
 PR 30-JAN-2001; 2001WO-US00668.
 PR 30-JAN-2001; 2001WO-US00669.
 PR 23-MAY-2001; 2001US-0864761.
 PR 09-OCT-2001; 2001US-0327898.
 XX
 PA (AEOM-) AEOMICA INC.
 XX
 PI Zhan J;
 XX
 DR WPI; 2002-676582/73.
 XX
 PT Novel isolated human testis expressed Patched like protein (HTPL),
 PT useful for identifying agonist and antagonist and specific binding
 PT partners, and for treating subjects having defects in HTPL -

XX PS Example 2; Page 290; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL has two isoforms, with a few single base pair differences between the two. One of the single base pair changes introduces a premature stop codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL shares an overall structure organisation with the Patched protein. The shared structural features strongly imply that HTPL plays a role similar to that of Patched, and is a potential tumour suppressor. HTPL is mapped to human chromosome 10p12.1. HTPL and its coding sequence are important in regulating male germ cell development, and the HTPL gene was useful for diagnosing a disorder caused by mutation in HTPL, and in therapy and manufacture of a medicament for treatment or prevention of such disorder associated with decreased expression or activity of human HTPL. Such disorders include disorders of testis, or adrenal, adult and foetal liver, bone marrow, brain, kidney, lung, placenta, prostate, skeletal muscle or colon function. HTPL proteins and nucleic acids are clinically useful diagnostic markers and potential therapeutic agents for male infertility and cancer. The present oligonucleotide was used in an example from the invention.

XX SQ Sequence 17 BP; 8 A; 1 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 11 CTTCTCTTTT 2

RESULT 23
ABV80481/C
ID ABV80481 standard; DNA; 17 BP.
XX AC ABV80481;
XX DT 03-JAN-2003 (first entry)
XX DE Human HTPL scanning oligonucleotide SEQ ID 1727.
XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
KW human testis expressed Patched like protein; testis; adrenal; liver;
KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
OS Homo sapiens.
XX PN EP1229046-A2.
XX PD 07-AUG-2002.
XX PF 28-JAN-2002; 2002EP-0001167.
XX PR 30-JAN-2001; 2001WO-US00663.
PR 30-JAN-2001; 2001WO-US00664.
PR 30-JAN-2001; 2001WO-US00665.
PR 30-JAN-2001; 2001WO-US00667.
PR 30-JAN-2001; 2001WO-US00668.
PR 30-JAN-2001; 2001WO-US00669.
PR 23-MAY-2001; 2001US-0864761.
PR 09-OCT-2001; 2001US-0327898.
XX PA (AEOM-) AEOMICA INC.
XX Zhan J;
XX PI WPI; 2002-676582/73.
XX DR Novel isolated human testis expressed Patched like protein (HTPL),
XX PT

PT useful for identifying agonist and antagonist and specific binding partners, and for treating subjects having defects in HTPL -

XX PS Example 2; Page 290; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL has two isoforms, with a few single base pair differences between the two. One of the single base pair changes introduces a premature stop codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL shares an overall structure organisation with the Patched protein. The shared structural features strongly imply that HTPL plays a role similar to that of Patched, and is a potential tumour suppressor. HTPL is mapped to human chromosome 10p12.1. HTPL and its coding sequence are important in regulating male germ cell development, and the HTPL gene was useful for diagnosing a disorder caused by mutation in HTPL, and in therapy and manufacture of a medicament for treatment or prevention of such disorder associated with decreased expression or activity of human HTPL. Such disorders include disorders of testis, or adrenal, adult and foetal liver, bone marrow, brain, kidney, lung, placenta, prostate, skeletal muscle or colon function. HTPL proteins and nucleic acids are clinically useful diagnostic markers and potential therapeutic agents for male infertility and cancer. The present oligonucleotide was used in an example from the invention.

XX SQ Sequence 17 BP; 8 A; 1 C; 5 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 10 CTTCTCTTTT 1

RESULT 24
ABS74761/C
ID ABS74761 standard; DNA; 17 BP.
XX AC ABS74761;
XX DT 24-DEC-2002 (first entry)
XX DE Human PAPP-Ea associated 17-mer SEQ ID 287.
XX KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
KW dysgenetic pregnancy; primer; ss.
OS Homo sapiens.
XX PN US2002102252-A1.
XX PD 01-AUG-2002.
XX PF 06-APR-2001; 2001US-0827998.
XX PR 26-MAY-2000; 2000US-207456P.
XX PA (GUYV/) GU Y.
PA (SHAN/) SHANNON M E.
XX PI Gu Y, Shannon ME;
XX DR WPI; 2002-697817/75.
XX PT New isolated nucleic acid encoding an isoform of human pregnancy associated plasma protein E, for preventing or aborting pregnancy -
XX PS Example 2; Page 113; 353pp; English.
XX CC This invention describes a novel isolated nucleic acid that encodes

CC one of three new isoforms of human pregnancy associated plasma protein E,
 CC hPAPP-E. The products of the invention have abortive and contraceptive
 CC activity and can be used for gene therapy or in a vaccine. The nucleic
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
 CC used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 CC antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.

XX SQ Sequence 17 BP; 10 A; 2 C; 4 G; 1 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
 DB 17 CTTCTCTTTT 8

RESULT 25
 ABS74762/c
 ID ABS74762 standard; DNA; 17 BP.

AC ABS74762;
 XX
 XX
 XX 24-DEC-2002 (first entry)
 XX Human PAPP-Ea associated 17-mer SEQ ID 288.

XX PAPP-E; human; pregnancy associated plasma protein E; abortive;
 XX contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
 XX dysgenetic pregnancy; primer; ss.

XX Homo sapiens.

XX US2002102252-A1.

XX 01-AUG-2002.

XX 06-APR-2001; 2001US-0827998.

XX 26-MAY-2000; 2000US-207456P.

XX (GUY/) GU Y.

XX (SHAN/) SHANNON M E.

XX Gu Y, Shannon ME;

XX WPI; 2002-697817/75.

XX New isolated nucleic acid encoding an isoform of human pregnancy
 XX associated plasma protein E, for preventing or aborting pregnancy -

XX Example 2; Page 113; 353pp; English.

XX This invention describes a novel isolated nucleic acid that encodes
 CC one of three new isoforms of human pregnancy associated plasma protein E,
 CC hPAPP-E. The products of the invention have abortive and contraceptive
 CC activity and can be used for gene therapy or in a vaccine. The nucleic
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
 CC used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 CC antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.

XX SQ Sequence 17 BP; 10 A; 2 C; 4 G; 1 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
 DB 16 CTTCTCTTTT 7

RESULT 26
 ABS74763/c
 ID ABS74763 standard; DNA; 17 BP.

AC ABS74763;
 XX
 XX 24-DEC-2002 (first entry)
 XX Human PAPP-Ea associated 17-mer SEQ ID 289.

XX PAPP-E; human; pregnancy associated plasma protein E; abortive;
 XX contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
 XX dysgenetic pregnancy; primer; ss.

XX Homo sapiens.

XX US2002102252-A1.

XX 01-AUG-2002.

XX 06-APR-2001; 2001US-0827998.

XX 26-MAY-2000; 2000US-207456P.

XX (GUY/) GU Y.

XX (SHAN/) SHANNON M E.

XX Gu Y, Shannon ME;

XX WPI; 2002-697817/75.

XX New isolated nucleic acid encoding an isoform of human pregnancy
 XX associated plasma protein E, for preventing or aborting pregnancy -

XX Example 2; Page 113; 353pp; English.

XX This invention describes a novel isolated nucleic acid that encodes
 CC one of three new isoforms of human pregnancy associated plasma protein E,
 CC hPAPP-E. The products of the invention have abortive and contraceptive
 CC activity and can be used for gene therapy or in a vaccine. The nucleic
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
 CC used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 CC antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.

XX SQ Sequence 17 BP; 9 A; 2 C; 4 G; 2 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
 DB 15 CTTCTCTTTT 6


```

RESULT 27
ABS74764/c
ID ABS74764 standard; DNA; 17 BP.
XX
XX AC ABS74764;
XX
XX DT 24-DEC-2002 (first entry)
XX
XX DE Human PAPP-Ea associated 17-mer SEQ ID 290.
XX
XX KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX KW dysgenetic pregnancy; primer; ss.
XX
XX OS Homo sapiens.
XX
XX PN US2002102252-A1.
XX
XX PD 01-AUG-2002.
XX
XX PF 06-APR-2001; 2001US-0827998.
XX
XX PR 26-MAY-2000; 2000US-207456P.
XX
XX PA (GUY/) GU Y.
XX PA (SHAN/) SHANNON M E.
XX
XX PI Gu Y, Shannon ME;
XX
XX DR WPI; 2002-697817/75.
XX
XX PT New isolated nucleic acid encoding an isoform of human pregnancy
XX PT associated plasma protein E, for preventing or aborting pregnancy -
XX
XX PS Example 2; Page 113; 353pp; English.
XX
XX CC This invention describes a novel isolated nucleic acid that encodes
XX CC one of three new isoforms of human pregnancy associated plasma protein E,
XX CC hPAPP-E. The products of the invention have abortive and contraceptive
XX CC activity and can be used for gene therapy or in a vaccine. The nucleic
XX CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
XX CC used in pharmaceutical compositions or vaccines for preventing or
XX CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
XX CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
XX CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
XX CC antibodies can be used to assess the expression levels of PAPP-E isoform
XX CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
XX CC antenatally. This sequence represents an oligomer used in scanning the
XX CC human PAPP-E genes described in the disclosure of the invention.
XX
XX SQ Sequence 17 BP; 10 A; 1 C; 4 G; 2 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
Best Local Similarity 100.0%; Pred. NO. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
DB 14 CTTCTCTTTT 5

RESULT 28
ABS74765/c
ID ABS74765 standard; DNA; 17 BP.
XX
XX AC ABS74765;
XX
XX DT 24-DEC-2002 (first entry)
XX
XX DE Human PAPP-Ea associated 17-mer SEQ ID 291.
XX
XX KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX KW

```

```

KW dysgenetic pregnancy; primer; ss.
XX
XX OS Homo sapiens.
XX
XX PN US2002102252-A1.
XX
XX PD 01-AUG-2002.
XX
XX PF 06-APR-2001; 2001US-0827998.
XX
XX PR 26-MAY-2000; 2000US-207456P.
XX
XX PA (GUY/) GU Y.
XX PA (SHAN/) SHANNON M E.
XX
XX PI Gu Y, Shannon ME;
XX
XX DR WPI; 2002-697817/75.
XX
XX PT New isolated nucleic acid encoding an isoform of human pregnancy
XX PT associated plasma protein E, for preventing or aborting pregnancy -
XX
XX PS Example 2; Page 113; 353pp; English.
XX
XX CC This invention describes a novel isolated nucleic acid that encodes
XX CC one of three new isoforms of human pregnancy associated plasma protein E,
XX CC hPAPP-E. The products of the invention have abortive and contraceptive
XX CC activity and can be used for gene therapy or in a vaccine. The nucleic
XX CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
XX CC used in pharmaceutical compositions or vaccines for preventing or
XX CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
XX CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
XX CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
XX CC antibodies can be used to assess the expression levels of PAPP-E isoform
XX CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
XX CC antenatally. This sequence represents an oligomer used in scanning the
XX CC human PAPP-E genes described in the disclosure of the invention.
XX
XX SQ Sequence 17 BP; 11 A; 1 C; 4 G; 1 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 17;
Best Local Similarity 100.0%; Pred. NO. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
DB 13 CTTCTCTTTT 4

RESULT 29
ABS74766/c
ID ABS74766 standard; DNA; 17 BP.
XX
XX AC ABS74766;
XX
XX DT 24-DEC-2002 (first entry)
XX
XX DE Human PAPP-Ea associated 17-mer SEQ ID 292.
XX
XX KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX KW dysgenetic pregnancy; primer; ss.
XX
XX OS Homo sapiens.
XX
XX PN US2002102252-A1.
XX
XX PD 01-AUG-2002.
XX
XX PF 06-APR-2001; 2001US-0827998.
XX
XX PR 26-MAY-2000; 2000US-207456P.
XX

```

PA (GUY/) GU Y.
 PA (SHAN/) SHANNON M E.
 XX
 PI Gu Y, Shannon ME;
 XX
 DR WPI; 2002-697817/75.
 XX
 XX
 PT New isolated nucleic acid encoding an isoform of human pregnancy
 associated plasma protein E, for preventing or aborting pregnancy -
 XX
 XX Example 2; Page 113; 353pp; English.
 XX
 CC This invention describes a novel isolated nucleic acid that encodes
 one of three new isoforms of human pregnancy associated plasma protein E,
 hPAPP-E. The products of the invention have abortive and contraceptive
 activity and can be used for gene therapy or in a vaccine. The nucleic
 acid, polypeptide encoded by it, or antibody to the polypeptide can be
 used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.
 XX
 SQ Sequence 17 BP; 10 A; 1 C; 4 G; 2 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTCTCTCTTT 10
 Db 12 CTCTCTCTTT 3
 RESULT 30
 ABS74767/C
 ID ABS74767 standard; DNA; 17 BP.
 XX
 AC ABS74767;
 XX
 DT 24-DEC-2002 (first entry)
 XX
 DE Human PAPP-Ea associated 17-mer SEQ ID 293.
 XX
 KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
 KW dysgenetic pregnancy; primer; ss.
 XX
 OS Homo sapiens.
 XX
 PN US2002102252-A1.
 XX
 PD 01-AUG-2002.
 XX
 PF 06-APR-2001; 2001US-0827998.
 XX
 PR 26-MAY-2000; 2000US-207456P.
 XX
 PA (GUY/) GU Y.
 PA (SHAN/) SHANNON M E.
 XX
 PI Gu Y, Shannon ME;
 XX
 DR WPI; 2002-697817/75.
 XX
 XX
 PT New isolated nucleic acid encoding an isoform of human pregnancy
 associated plasma protein E, for preventing or aborting pregnancy -
 XX
 XX Example 2; Page 113; 353pp; English.
 XX

CC This invention describes a novel isolated nucleic acid that encodes
 one of three new isoforms of human pregnancy associated plasma protein E,
 hPAPP-E. The products of the invention have abortive and contraceptive
 activity and can be used for gene therapy or in a vaccine. The nucleic
 acid, polypeptide encoded by it, or antibody to the polypeptide can be
 used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.
 XX
 SQ Sequence 17 BP; 10 A; 0 C; 4 G; 3 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTCTCTCTTT 10
 Db 11 CTCTCTCTTT 2
 RESULT 31
 ABS74768/C
 ID ABS74768 standard; DNA; 17 BP.
 XX
 AC ABS74768;
 XX
 DT 24-DEC-2002 (first entry)
 XX
 DE Human PAPP-Ea associated 17-mer SEQ ID 294.
 XX
 KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
 KW dysgenetic pregnancy; primer; ss.
 XX
 OS Homo sapiens.
 XX
 PN US2002102252-A1.
 XX
 PD 01-AUG-2002.
 XX
 PF 06-APR-2001; 2001US-0827998.
 XX
 PR 26-MAY-2000; 2000US-207456P.
 XX
 PA (GUY/) GU Y.
 PA (SHAN/) SHANNON M E.
 XX
 PI Gu Y, Shannon ME;
 XX
 DR WPI; 2002-697817/75.
 XX
 XX
 PT New isolated nucleic acid encoding an isoform of human pregnancy
 associated plasma protein E, for preventing or aborting pregnancy -
 XX
 XX Example 2; Page 113; 353pp; English.
 XX

CC This invention describes a novel isolated nucleic acid that encodes
 one of three new isoforms of human pregnancy associated plasma protein E,
 hPAPP-E. The products of the invention have abortive and contraceptive
 activity and can be used for gene therapy or in a vaccine. The nucleic
 acid, polypeptide encoded by it, or antibody to the polypeptide can be
 used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.
 XX
 SQ Sequence 17 BP; 10 A; 0 C; 4 G; 3 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 17;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTCTCTCTTT 10
 Db 11 CTCTCTCTTT 2
 RESULT 31
 ABS74768/C
 ID ABS74768 standard; DNA; 17 BP.
 XX
 AC ABS74768;
 XX
 DT 24-DEC-2002 (first entry)
 XX
 DE Human PAPP-Ea associated 17-mer SEQ ID 294.
 XX
 KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
 KW dysgenetic pregnancy; primer; ss.
 XX
 OS Homo sapiens.
 XX
 PN US2002102252-A1.
 XX
 PD 01-AUG-2002.
 XX
 PF 06-APR-2001; 2001US-0827998.
 XX
 PR 26-MAY-2000; 2000US-207456P.
 XX
 PA (GUY/) GU Y.
 PA (SHAN/) SHANNON M E.
 XX
 PI Gu Y, Shannon ME;
 XX
 DR WPI; 2002-697817/75.
 XX
 XX
 PT New isolated nucleic acid encoding an isoform of human pregnancy
 associated plasma protein E, for preventing or aborting pregnancy -
 XX
 XX Example 2; Page 113; 353pp; English.
 XX

CC human PAPP-E genes described in the disclosure of the invention.

XX Sequence 17 BP; 9 A; 1 C; 4 G; 3 T; 0 other;
SQ Query Match 100.0%; Score 10; DB 24; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 10 CTTCTCTTTT 1

RESULT 32
ABX94543
ID ABX94543 standard; DNA; 19 BP.

XX AC ABX94543;
XX 13-JUN-2003 (first entry)
XX 23S/16S rRNA detecting probe SEQ ID 12.
XX Detection; probe; contaminant; drinking water; Legionella; coliform;
XX faecal streptococci; soil; sputum; biopsy; urine; food; pharmaceutical;
XX cosmetic; fluorescent in situ hybridisation; FISH; ss.
XX Streptococcus sp.
XX WO2002102824-A2.
XX 27-DEC-2002.

XX 19-JUN-2002; 2002WO-EP06809.
XX 19-JUN-2001; 2001DE-1029411.
XX 11-DEC-2001; 2001DE-1060666.

XX (VERM-) VERMICON AG.

PI Beinfuhr C, Snaidr J;

XX WPI; 2003-167479/16.

XX New oligonucleotides, useful for detecting bacteria that may
PT contaminate drinking water, provide quick results for many species in
PT parallel -

PS Claim 8; Page 12; 53pp; German.

XX This invention describes novel oligonucleotide probes used to detect
CC contaminant bacteria that may be present in drinking water. The probes
CC can detect bacteria (especially Legionella, faecal streptococci and
CC coliforms) that may contaminate drinking water in environmental samples
CC (water or soil), clinical samples (sputum, biopsies, urine etc.), in
CC bathing and drinking water and in foods, pharmaceuticals and cosmetics,
CC by in situ hybridisation. The probes combine the advantages of
CC fluorescent in situ hybridisation with those of culture methods. Only a
CC relatively short culture step is required; analysis takes 24-48 hours
CC (contrast many days for conventional methods) and all relevant bacteria
CC can be tested simultaneously. The oligonucleotides can differentiate
CC between species of the same genus and are easy to use, allowing simple
CC analysis of a large number of samples. ABX94532-ABX94578 represent the
CC oligonucleotide probes described in the invention.

XX Sequence 19 BP; 1 A; 7 C; 2 G; 9 T; 0 other;

Query Match 100.0%; Score 10; DB 25; Length 19;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||

Db 4 CTTCTCTTTT 13

RESULT 33

AA76779
ID AA76779 standard; DNA; 20 BP.

XX AC AA76779;

XX 15-SEP-1997 (first entry)

XX Staphylococcus aureus exfoliative toxin A gene PCR primer ETA-B.

XX Asymmetric polymerase chain reaction; nucleic acid amplification;
KW PCR; detection; assay; exfoliative toxin A; ETA; skin lesion;
KW competitive primer; capture probe; ss.

OS Synthetic.

XX US5627054-A.

XX 06-MAY-1997.

XX 05-APR-1996; 96US-0628417.

XX 05-APR-1996; 96US-0628417.

XX (USSA) US SEC OF ARMY.

XX Gillespie D;

XX WPI; 1997-271311/24.

XX Quantitative nucleic acid amplification - by competitor primer
PT asymmetric polymerase chain reaction

XX Example 1; Column 5; 9pp; English.

CC In a specific example of a novel process for amplifying an amount
CC (known or unknown) of a double-stranded nucleic acid segment to produce
CC single-stranded nucleic acid in an amount that is proportional to the
CC starting amount of the nucleic acid, the Staphylococcus aureus
CC exfoliative toxin A (ETA) gene was used as the DNA template. The
CC region comprising nucleotides 165-436 was amplified by symmetric,
CC asymmetric or competitor primer asymmetric PCR using the primers
CC ETA-A2 and ETA-B (see AA76778 and AA76779). For asymmetric PCR, the
CC amount of primer ETA-B was reduced and for competitor primer
CC asymmetric PCR a competitor primer ETA-B2 (see AA76780) was added
CC with upstream primer ETA-A2 after the initial cycling reaction. PCR
CC products containing ETA-specific sequences were detected
CC radioactively by a capture system which employed a bifunctional
CC capture probe ETA-CP (see AA76781 and AA76782). ETA-CP was designed
CC to capture the amplified sense strand onto capture membranes
CC through hybridisation between the first 40 nucleotides of ETA-CP
CC and nucleotides 321-360 of the ETA gene and through hybridisation
CC of the poly(dA) tail on ETA-CP with poly(dT) tails on the capture
CC membranes. A radioactively labelled "label probe" (see AA76783),
CC complementary to nucleotides 389-410 of the ETA gene was used to
CC detect the amplicons. Results showed that hybridisation of the
CC capture probe and label probe to the denatured symmetric PCR
CC product was much less efficient than hybridisation to the
CC single-stranded PCR products of the asymmetric and competitor
CC primer asymmetric reactions.

XX Sequence 20 BP; 4 A; 5 C; 3 G; 8 T; 0 other;

Query Match 100.0%; Score 10; DB 18; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 8 CTTCTCTTTT 17

RESULT 34
AAZ04007
ID AAZ04007 standard; DNA; 20 BP.
XX AC
XX AAZ04007;
XX
DT 07-OCT-1999 (first entry)
XX
DE PCR primer used to amplify an ORF of Chlamydia trachomatis.
XX
KW Vaccine; eye disease; conventional trachoma; nonendemic trachoma;
KW paratrachoma; inclusion conjunctivitis; genital disease; perihhepatitis;
KW nongonococcal urethritis; epididymitis; cervicitis; salpingitis; PCR primer;
KW Bartholinitis; pneumopathy; venereal lymphogranulomatosis; ss.
XX
OS Synthetic.
OS Chlamydia trachomatis.
XX
PN WO9928475-A2.
XX
PD 10-JUN-1999.
XX
XX 27-NOV-1998; 98WO-IB01939.
XX
PR 04-NOV-1998; 98US-0107077.
PR 28-NOV-1997; 97FR-0015041.
PR 17-DEC-1997; 97FR-0016034.
XX
XX (GEST) GENSET.
XX
PA Griffais R;
PI
XX WPI; 1999-371125/31.
XX
XX Genome sequence of Chlamydia trachomatis
PT
XX Disclosure; Page 1653; 1755pp; English.
XX
XX PCR primers AAZ01426-Z06209 were used to amplify open reading frames
CC (ORFs) of the genome of Chlamydia trachomatis (see AAZ01425). These ORFs
CC encode polypeptides (see AAX36754-Y37949) which can be used as vaccines
CC against Chlamydia trachomatis. Antisense and ribozyme sequences
CC can also be used to control growth of the microorganism. Chlamydia
CC trachomatis is responsible for a large number of diseases, e.g. eye
CC diseases such as conventional trachoma, nonendemic trachoma,
CC paratrachoma, and inclusion conjunctivitis; genital diseases such as
CC nongonococcal urethritis, epididymitis, cervicitis, salpingitis,
CC perihhepatitis, Bartholinitis; pneumopathy in breast feeding infants;
CC and venereal lymphogranulomatosis. The polypeptides of the
CC invention may be of use in treating these diseases.
XX
SQ Sequence 20 BP; 0 A; 7 C; 3 G; 10 T; 0 other;

Query Match 100.0%; Score 10; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 3 CTTCTCTTTT 12

RESULT 35
AAX97056/c
ID AAX97056 standard; DNA; 20 BP.
XX AC
XX AAX97056;
XX
DT 13-SEP-1999 (first entry)
XX
DE PCR primer used to amplify an ORF of Chlamydia pneumoniae.

XX Respiratory disease; pneumonia; bronchitis; heart disease; sarcoidosis;
KW sinusitis; purulent otitis media; erythema nodosum; pharyngitis;
KW vaccine; neutralising epitope; PCR primer; ss.
XX
OS Synthetic.
OS Chlamydia pneumoniae.
XX
PN WO9927105-A2.
XX
XX 03-JUN-1999.
XX
PD 20-NOV-1998; 98WO-IB01890.
XX
PF 04-NOV-1998; 98US-0107078.
PR 21-NOV-1997; 97FR-0014673.
XX
XX (GEST) GENSET.
XX
PA Griffais R;
PI
XX WPI; 1999-357842/30.
XX
XX Genome sequence of Chlamydia pneumoniae
PT
XX Page 1874; Disclosure; 1912pp; English.
XX
XX AAX91991-X97517 represent PCR primers used to amplify open reading
CC frames and other nucleic acid sequences from the genome of
CC Chlamydia pneumoniae (see AAX91990). C. pneumoniae causes respiratory
CC disease such as pneumonia and bronchitis and is thought to be a
CC contributing factor in heart disease, sarcoidosis, sinusitis, purulent
CC otitis media, erythema nodosum or pharyngitis. The polypeptides encoded
CC by the open reading frames of the C. pneumoniae genome (see AAX94584-
CC AAX35879) can be used in immunogenic compositions as vaccines. Vectors
CC containing C. pneumoniae nucleotides sequences can also be used as
CC immunogenic compositions, especially where the vector directs the
CC expression of a neutralising epitope of C. pneumoniae.
XX
SQ Sequence 20 BP; 8 A; 4 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 20; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 17 CTTCTCTTTT 8

RESULT 36
AAL51617/c
ID AAL51617 standard; DNA; 20 BP.
XX AC
XX AAL51617;
XX
DT 17-APR-2003 (first entry)
XX
XX Human interferon alpha 2 sequencing primer #5.
XX
XX Human; ss; cellular proliferation inhibitor; interferon alpha 2;
KW single nucleotide polymorphism; SNP; cancer; tumour; metabolic disease;
KW cardiovascular disease; infectious disease; immunological disease; HIV;
KW central nervous system disease; wound healing; chemotherapy side effect;
KW anaemia; osteoporosis; gastrointestinal disease; venereal disease; AIDS;
KW obesity; hepatitis; infectious pneumonia; Alzheimer's disease; allergy;
KW Parkinson's disease; multiple sclerosis; schizophrenia; depression;
KW graft versus host disease; asthma; psoriasis; rheumatoid arthritis;
KW Crohn's disease; ulcerative colitis; genital wart; sequencing; primer.
XX
OS Homo sapiens.
XX
PN EP1236800-A2.

XX PD 04-SEP-2002.
 XX PF
 XX PR 01-MAR-2002; 2002EP-0290515.
 XX PR 01-MAR-2001; 2001FR-0002843.
 XX PA (GENO-) GENODYSSEE.
 XX PI Escary J;
 XX PI WPI; 2003-185789/19.
 XX DR
 XX PT An isolated polynucleotide encoding interferon alpha 2 containing
 PT single nucleotide polymorphisms is useful in treating disease -
 XX
 XX PS Example 4; Page 21; 42pp; English.
 XX CC The invention comprises the amino acid and coding sequence of the human
 CC interferon alpha 2 protein. The invention further relates to the
 CC identification of single nucleotide polymorphisms (SNPs) within the human
 CC interferon alpha 2 gene. The DNA and protein sequences of the invention
 CC are useful for the treatment of: cancer; tumours; cardiovascular
 CC diseases; metabolic diseases; infectious diseases; central nervous system
 CC diseases; immunological diseases; wound healing; chemotherapy side
 CC effects; anaemia; osteoporosis; Gastrointestinal diseases; venereal
 CC diseases; obesity; hepatitis; HIV/AIDS; infectious pneumonias;
 CC Alzheimer's disease; Parkinson's disease; multiple sclerosis;
 CC schizophrenia; depression; graft versus host disease; allergies; asthma;
 CC psoriasis; rheumatoid arthritis; Crohn's disease; ulcerative colitis; and
 CC genital warts. The present DNA sequence represents a primer that was used
 CC to sequence the human interferon alpha 2 gene.
 XX
 XX SQ Sequence 20 BP; 14 A; 1 C; 5 G; 0 U; 0 other;
 Query Match 100.0%; Score 10; DB 25; Length 20;
 Best Local Similarity 100.0%; Pred No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 |||||
 Db 20 CTTCTCTTTT 11
 RESULT 37
 AAX60140/c
 ID AAX60140 standard; DNA; 21 BP.
 XX AC
 XX AAX60140;
 XX DT 05-AUG-1999 (first entry)
 XX DE PCR primer used to amplify Mycoplasma hyopneumoniae P102 protein DNA.
 XX KW P102 protein; vaccine; antigen; diagnosis; swine; immunisation;
 KW enzootic pneumonia; PCR primer; ss.
 XX OS Synthetic.
 XX PN WO926664-A1.
 XX PD 03-JUN-1999.
 XX PF 24-NOV-1998; 98WO-US25044.
 XX PR 26-NOV-1997; 97US-0066565.
 XX PA (IOWA) UNIV IOWA STATE RES FOUND INC.
 XX PI Hsu T, Minion PC;
 XX DR WPI; 1999-357741/30.
 XX

PT Recombinant antigenic Mycoplasma hyopneumoniae protein
 XX
 XX PS Example 2; Page 23; 45pp; English.
 XX CC PCR primers AAX60140-41 were used to amplify DNA encoding a Mycoplasma
 CC hyopneumoniae P102 protein clone. The P102 protein and its fragments
 CC are used in vaccines to protect against enzootic pneumonia,
 CC particularly in swine. Recombinant P102 polypeptides may be used as
 CC antigens for diagnostic purposes to determine whether or not a
 CC biological test sample contains M. hyopneumoniae antigens or
 CC antibodies. The P102 polypeptides or DNA sequences may also be used
 CC for immunising or protecting non-human animals, preferably swine,
 CC against M. hyopneumoniae infections, particularly enzootic pneumonia.
 XX
 XX SQ Sequence 21 BP; 12 A; 0 C; 7 G; 2 T; 0 other;
 Query Match 100.0%; Score 10; DB 20; Length 21;
 Best Local Similarity 100.0%; Pred No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 |||||
 Db 15 CTTCTCTTTT 6
 RESULT 38
 AAV08126/c
 ID AAV08126 standard; DNA; 21 BP.
 XX AC AAV08126;
 XX DT 22-JAN-1999 (first entry)
 XX DE Primer Vbeta14 for T cell receptor V region.
 XX KW PCR primer; T-cell receptor; TCR; V region; immune response; arthritis;
 KW somatic homologous recombination; hypervariable region;
 KW spectratype determination; autoimmune response; multiple sclerosis;
 KW myasthenia gravis; muscular dystrophy; graft-infiltrating lymphocyte;
 KW tumour-infiltrating lymphocyte; ss.
 XX OS Synthetic.
 XX OS Mammalia.
 XX PN US5837447-A.
 XX PD 17-NOV-1998.
 XX PF 19-APR-1994; 94US-0229528.
 XX PR 19-APR-1994; 94US-0229528.
 XX PR 15-APR-1992; 92US-0868569.
 XX PA (BLOO-) BLOOD CENT RES FOUND INC.
 XX PI Gorski J;
 XX DR WPI; 1999-023435/02.
 XX PT Monitoring immune responses by analysing amplified B or T-cell
 PT nucleic acid - using primers specific for variable and constant or
 PT junction region gene segments, with separation of products by
 PT length, especially to monitor auto:immunity
 XX
 XX PS Claim 22; Column 38; 26pp; English.
 XX CC This sequence represents a primer for the T cell receptor (TCR) variable
 CC region and is used in the method of the invention. The method is for
 CC monitoring an immune response that involves somatic homologous
 CC recombination between elements of at least two segments associated with a
 CC hypervariable region, and comprises: (a) providing a polynucleotide
 CC sample from B- or T-cells, and amplifying it with: (i) a primer specific
 CC for a variable gene segment; and (ii) a primer specific for a constant or

CC joining gene segment to produce amplification products (AP) that can be
 CC resolved at a difference in size of 2 or 3 bp; (c) separating the AP
 CC according to length; (d) detecting the range of lengths in the separated
 CC products to produce a 'spectratype' of the subject's immune response; and
 CC (e) comparing the spectratype with a predetermined standard to determine
 CC immune status or to monitor immune response. The method is specifically
 CC used to monitor autoimmune responses (including relapses), i.e. to
 CC identify the predominant TCR in sites of autoimmune activity (e.g. in
 CC arthritis, multiple sclerosis, myasthenia gravis and muscular dystrophy)
 CC or present in graft-infiltrating (in cases of organ rejection) or
 CC tumour-infiltrating lymphocytes. As each gene rearrangement is unique,
 CC each complementarity determining region 3 is a specific molecular
 CC fingerprint of the lymphocyte that generates it, and immune responses can
 CC be correlated with an increase in a particular TCR or immunoglobulin.
 CC Specific determination of two V beta families may be done simultaneously.
 XX Sequence 21 BP; 8 A; 3 C; 7 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 20; Length 21;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
 Db 18 CTTCTCTTTT 9

RESULT 39
 AAA95613/c
 ID AAA95613 standard; DNA; 21 BP.

AC AAA95613;
 DT 31-JAN-2001 (first entry)
 DE TCR Vbeta 14 subfamily probe VB14-1.
 XX
 XX Detection; diagnostic; Kawasaki disease; T-cell; PCR primer; probe;
 KW gene expression; ss.
 XX Homo sapiens.

PN JP2000157297-A.
 XX 13-JUN-2000.
 PD 01-DEC-1998; 98JP-0341661.
 XX 01-DEC-1998; 98JP-0341661.
 PR (SHIO) SHIONOGI & CO LTD.
 XX WPI; 2000-477722/42.
 DR
 XX Detection of Kawasaki disease factor, useful for the diagnosis of
 PT Kawasaki disease, comprises detecting an increase in Vbeta6.5 positive
 PT T-cells -
 XX

PS Example 1; Page 9; 36pp; Japanese.

XX The invention relates to a method of detecting Kawasaki disease by
 CC detecting an increase in Vbeta6.5 or Vbeta6.5/Vbeta2.1 positive T-cells.
 CC The sequences AAA95613-A95626 represent primers and probes used to PCR
 CC amplify and detect the level of expression of Valpha and Vbeta genes
 CC in T-cells in Kawasaki disease.

XX Sequence 21 BP; 8 A; 3 C; 7 G; 3 T; 0 other;

Query Match 100.0%; Score 10; DB 21; Length 21;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

|||||

18 CTTCTCTTTT 9

RESULT 40
 AAH27157/c
 ID AAH27157 standard; DNA; 21 BP.

XX AAH27157;
 AC
 XX 08-AUG-2001 (first entry)
 DT
 XX Downstream PCR primer for amplification of PDAT gene.
 DE
 XX Yeast; ARE 1; YCR048w; transgenic plant; oil production; acyl-CoA;
 KW fatty acid production; triacylglycerol; oil crop; rape; sunflower; PDAT;
 KW oil palm; soy; maize; oat; potato; sugar beet; turnip; PCR primer; ss.
 XX

XX Saccharomyces cerevisiae.

XX WO200134814-A1.

XX 17-MAY-2001.

XX 10-NOV-2000; 2000WO-SB02216.

XX 12-NOV-1999; 93EP-0850169.

XX 12-NOV-1999; 99US-0164859.

XX (SCAN-) SCANBI SCANDINAVIAN BIOTECHNOLOGY RES AB.

XX Banas A, Sandager L, Stahl U, Dahlqvist A, Lenman M, Ronne H;
 PI Stymne S;

XX WPI; 2001-329086/34.

XX Transforming oil-producing organisms with a gene encoding an
 PT acyl-CoA:diacylglycerol acyltransferase, useful to generate
 PT agricultural crops with higher triacylglycerol content -

PS Example 1; Page 6; 30pp; English.

XX This invention relates to the use of a novel enzyme in the production of
 CC an oil-producing organism. The enzyme catalyses the transfer of a fatty
 CC acid from acyl-CoA to diacylglycerol to produce triacylglycerol,
 CC resulting in an increased oil content. Sequences AAH27155 and AAH97263
 CC represent the Saccharomyces cerevisiae ARE1 coding and protein sequence
 CC respectively. ARE1 is used in the transformation or the organism of the
 CC invention. The invention is used to increase the oil content of oil crops
 CC such as rape, sunflower and oil palm, and other crops such as soy, maize,
 CC oat, potato, sugar beet and turnips. The invention could also be used to
 CC produce triacylglycerols in microorganisms. The present sequence
 CC represents a PCR primer used to amplify the Saccharomyces cerevisiae PDAT
 CC gene. The primer and PCR product are used in an example illustrating that
 CC triacylglycerol accumulation is reduced in yeast cells that lack the ARE1
 CC gene. The primer is specifically used in the production of mutant yeast
 CC strains.

XX Sequence 21 BP; 13 A; 3 C; 5 G; 0 U; 0 other;

Query Match 100.0%; Score 10; DB 22; Length 21;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

|||||

17 CTTCTCTTTT 8

RESULT 41
 AAS95067/c
 ID AAS95067 standard; DNA; 21 BP.

XX

```

AC AAS95067;
DT 13-FEB-2002 (first entry)
DE Human otoferlin exon PCR primer #32.
KW Human; mouse; otoferlin; OTOF; brain; auditory function; PCR primer;
KW autosomal nonsyndromic prelingual deafness; DFNB9; ss.
OS Homo sapiens.
PN WO200170972-A2.
XX 27-SEP-2001.
XX 23-MAR-2001; 2001WO-IB00578.
XX 24-MAR-2000; 2000US-191738P.
XX (INSP ) INST PASTEUR.
XX PA (CNRS ) CNRS CENT NAT RECH SCI.
XX Yasnaga S, Grati M, Cohen-Salmon M, El Amraoui A, Petit C;
XX Weil D;
XX WPI; 2001-611499/70.
XX Novel human gene Otoferlin, underlying an autosomal recessive
XX nonsyndromic prelingual deafness, DFNB9, and proteins encoded by the
XX gene, implicated in deafness -
XX Claim 25; Page 17; 99pp; English.
XX
XX The invention relates to a purified polynucleotide (I) encoding a protein
XX sequence (II) encoded by a novel human gene, otoferlin (OTOF) or
XX the long human otoferlin isoform in brain. (I) was identified as
XX underlying an autosomal nonsyndromic prelingual deafness DFNB9, and is
XX thus useful for detecting deafness disease in humans and for
XX characterising the functions of proteins and genes encoding them in
XX auditory function. AAS95022-AAS95248 represent human and mouse
XX otoferlin coding sequences, PCR primers and related sequences of the
XX invention.
XX
XX Sequence 21 BP; 10 A; 1 C; 8 G; 2 T; 0 other;
XX
Query Match 100.0%; Score 10; DB 23; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
DB 10 CTTCTCTTTT 1
RESULT 42
AAX14678
ID AAX14678 standard; DNA; 22 BP.
XX
XX AAX14678;
XX
XX 24-MAR-1999 (first entry)
XX
XX Triple helix forming nucleotides 9-30 of gamma-crystallin gene.
XX
XX Triple-helix forming region; Triplex formation; DNA detection;
XX identification; bacteria; oncogene; virus; ds.
XX
XX Homo sapiens.
XX
XX US5861244-A.
XX
XX 19-JAN-1999.
XX

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PF 22-DEC-1993; 93US-0173489.
XX
XX 22-DEC-1993; 93US-0173489.
PR 29-OCT-1992; 92US-0968436.
XX
XX (PROF-) PROFILE DIAGNOSTIC SCI INC.
XX Hepburn AG, Wang C;
PI
XX WPI; 1999-130384/11.
XX
XX Assay of genetic sequences based on triplex formation from double
XX stranded analyte - and hybrid of anchor and reporter sequences, with
XX reporter released if triplex formation occurs, used e.g. to identify
XX bacteria
XX
XX Disclosure; Columns 15-16; 168pp; English.
XX
XX The present sequence represents a potential triple-helix forming region.
XX It can be used to demonstrate the assay of the invention. The assay
XX comprises adding a sample containing double-stranded DNA test sequences,
XX e.g. containing the present sequence, to an aqueous medium containing at
XX least one complex of anchor DNA, attached to a solid support, and
XX reporter DNA, where either a part of the anchor DNA or reporter DNA is
XX designed to form a triple-strand structure with part of the test
XX sequence. Triplex formation results in displacement of the reporter DNA
XX which is detected as an indication of the presence of the DNA test
XX sequence. The method is used to detect DNA sequences, particularly for
XX identification of bacteria (by detecting genes for ribosomal RNA) in
XX clinical samples, but also detection of oncogenes and Hepatitis B virus.
XX
XX Sequence 22 BP; 1 A; 5 C; 0 G; 16 T; 0 other;
XX
Query Match 100.0%; Score 10; DB 20; Length 22;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
DB 6 CTTCTCTTTT 15
RESULT 43
AAC66394/c
ID AAC66394 standard; DNA; 22 BP.
XX
XX AAC66394;
XX
XX 26-FEB-2001 (first entry)
XX
XX Human 3-hydroxyacyl-CoA-dehydratase cDNA specific PCR primer.
XX
XX Human; 3-hydroxyacyl-CoA-dehydratase; HCDase; hypothalamus;
XX PCR primer; ss.
XX
XX Homo sapiens.
XX
XX CN1263157-A.
XX
XX 16-AUG-2000.
XX
XX 17-FEB-2000; 2000CN-0111690.
XX
XX 17-FEB-2000; 2000CN-0111690.
XX
XX (NANF-) NANFANG RES CENT STATE HUMAN GENE GROUP.
XX
XX Li N, Qian B, Peng Y;
XX
XX WPI; 2000-639262/62.
XX
XX New human hydroxybutyryl coenzyme A dehydratase protein and its coding
XX sequence -

```

XX Example 1; Page 11; 21pp; Chinese.

PS This invention relates to a new human 3-hydroxyacyl-CoA-dehydratase

XX (HCDase). The protein is expressed in normal hypothalamic tissue in

CC humans. The invention includes human HCDase nucleotide and amino acid

CC sequences, a method for the preparation of the protein and a method for

CC detecting human HCDase nucleic acid and protein sequences in a sample.

CC The present sequence represents a PCR primer specific for human HCDase

CC cDNA.

XX

XX Sequence 22 BP; 10 A; 2 C; 8 G; 2 T; 0 other;

Query Match 100.0%; Score 10; DB 21; Length 22;

Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

Db 16 CTTCTCTTTT 7

RESULT 44

ABL35686

ID ABL35686 standard; DNA; 22 BP.

XX

AC ABL35686;

XX

DT 04-APR-2002 (first entry)

XX

DE Immunostimulatory oligonucleotide SEQ ID NO: 612.

XX

KW DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory;

KW vaccine; infection; allergy; cancer; hypersensitivity; bio-warfare;

KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;

KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;

KW antiinflammatory; antibacterial; ss.

XX

OS Synthetic.

XX

FH Key Location/Qualifiers

FT misc_RNA 1..22

FT /tag= a

FT /note= "optionally thymidine is replaced by uracil to

FT form RNA or DNA/RNA hybrids. Thymidine is linked to at

FT least one other base through a ribose sugar"

XX

PN WO200193902-A2.

XX

PD 13-DEC-2001.

XX

PF 07-JUN-2001; 2001WO-US18276.

XX

PR 07-JUN-2000; 2000US-209797P.

XX

PA (BIOS-) BIOSYNEXUS INC.

XX

PI Mond JJ, Flora M, Klinman DM;

XX

DR WPI; 2002-130570/17.

XX

PT New immunostimulatory compositions comprising RNA/DNA hybrid

PT oligonucleotides, useful for enhancing an immune response or inducing

PT cytokines, particularly for treating diseases, e.g. cancer, allergy or

PT HIV infection -

XX

PS Example 11; Page 63; 68pp; English.

XX

CC The present invention relates to an immunostimulatory composition, which

CC comprises at least one oligonucleotide comprising both an RNA region and

CC a DNA region. The composition is useful for enhancing an immune response

CC or inducing cytokines. It can be used as a vaccine adjuvant and in

CC treating diseases, including pathogenic infection, (non-)malignant

CC tumors (e.g. cancers of the brain, lung, ovary, breast, prostate or

CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies

CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or

CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies

CC (e.g. allergic rhinitis, hay fever, or food allergies), Lyme disease,

CC hepatitis, HIV or malaria. The composition is also useful for treating,

CC preventing or ameliorating the symptoms resulting from exposure to a

CC bio-warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence

CC is an immunostimulatory oligonucleotide described in the exemplification

XX of the invention.

XX Sequence 22 BP; 1 A; 4 C; 3 G; 14 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 22;

Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

Db 7 CTTCTCTTTT 16

RESULT 45

ABL35687

ID ABL35687 standard; DNA; 22 BP.

XX

AC ABL35687;

XX

DT 04-APR-2002 (first entry)

XX

DE Immunostimulatory oligonucleotide SEQ ID NO: 613.

XX

KW DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory;

KW vaccine; infection; allergy; cancer; hypersensitivity; bio-warfare;

KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;

KW immunosuppressive; protozoacide; virucide; hepatotropic; gene therapy;

KW antiinflammatory; antibacterial; ss.

XX

OS Synthetic.

XX

FH Key Location/Qualifiers

FT misc_RNA 1..22

FT /tag= a

FT /note= "optionally thymidine is replaced by uracil to

FT form RNA or DNA/RNA hybrids. Thymidine is linked to at

FT least one other base through a ribose sugar"

XX

PN WO200193902-A2.

XX

PD 13-DEC-2001.

XX

PF 07-JUN-2001; 2001WO-US18276.

XX

PR 07-JUN-2000; 2000US-209797P.

XX

PA (BIOS-) BIOSYNEXUS INC.

XX

PI Mond JJ, Flora M, Klinman DM;

XX

DR WPI; 2002-130570/17.

XX

PT New immunostimulatory compositions comprising RNA/DNA hybrid

PT oligonucleotides, useful for enhancing an immune response or inducing

PT cytokines, particularly for treating diseases, e.g. cancer, allergy or

PT HIV infection -

XX

PS Example 11; Page 63; 68pp; English.

XX

CC The present invention relates to an immunostimulatory composition, which

CC comprises at least one oligonucleotide comprising both an RNA region and

CC a DNA region. The composition is useful for enhancing an immune response

CC or inducing cytokines. It can be used as a vaccine adjuvant and in

CC treating diseases, including pathogenic infection, (non-)malignant

CC tumors (e.g. cancers of the brain, lung, ovary, breast, prostate or

CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies

CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
 CC hepatitis, HIV or malaria. The composition is also useful for treating,
 CC preventing or ameliorating the symptoms resulting from exposure to a
 CC bio-warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence
 CC is an immunostimulatory oligonucleotide described in the exemplification
 CC of the invention.

XX Sequence 22 BP; 1 A; 4 C; 4 G; 13 T; 0 other;

Qy Query Match 100.0%; Score 10; DB 24; Length 22;
 Db Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 Db 7 CTTCTCTTTT 16

RESULT 46
 ABL35688
 ID ABL35688 standard; DNA; 22 BP.
 XX AC ABL35688;
 XX DT 04-APR-2002 (first entry)
 XX DE Immunostimulatory oligonucleotide SEQ ID NO: 614.
 XX KW DNA/RNA hybrid; phosphorothioate backbone; immunostimulatory;
 KW vaccine; infection; allergy; cancer; hypersensitivity; bio-warfare;
 KW immunostimulant; antiallergic; cytostatic; antimicrobial; anti-HIV;
 KW immunosuppressive; protozoicide; virucide; hepatotropic; gene therapy;
 KW antiinflammatory; antibacterial; ss.
 XX OS Synthetic.
 XX FH Key Location/Qualifiers
 FT misc_RNA 1..22
 FT /tag= a
 FT /note= "optionally thymidine is replaced by uracil to
 FT form RNA or DNA/RNA hybrids. Thymidine is linked to at
 FT least one other base through a ribose sugar"
 XX WO200193902-A2.
 XX PD 13-DEC-2001.
 XX PF 07-JUN-2001; 2001WO-US18276.
 XX PR 07-JUN-2000; 2000US-209797P.
 XX PA (BIOS-) BIOSYNEXUS INC.
 XX PI Mond JJ, Flora M, Klinman DM;
 XX WI; 2002-130570/17.
 XX New immunostimulatory compositions comprising RNA/DNA hybrid
 PT oligonucleotides, useful for enhancing an immune response or inducing
 PT cytokines, particularly for treating diseases, e.g. cancer, allergy or
 PT HIV infection -
 XX Example 11; Page 63; 68pp; English.

CC The present invention relates to an immunostimulatory composition, which
 CC comprises at least one oligonucleotide comprising both an RNA region and
 CC a DNA region. The composition is useful for enhancing an immune response
 CC or inducing cytokines. It can be used as a vaccine adjuvant and in
 CC treating diseases, including pathogenic infection, (non-)malignant
 CC tumours (e.g. cancers of the brain, lung, ovary, breast, prostate or
 CC colon, or carcinomas and sarcomas), autoimmune diseases or allergies
 CC (e.g. allergic rhinitis, hay fever or food allergies), Lyme disease,
 CC hepatitis, HIV or malaria. The composition is also useful for treating,

CC preventing or ameliorating the symptoms resulting from exposure to a
 CC bio-warfare agent, e.g. Ebola, Anthrax or Listeria. The present sequence
 CC is an immunostimulatory oligonucleotide described in the exemplification
 CC of the invention.

XX Sequence 22 BP; 1 A; 4 C; 2 G; 15 T; 0 other;

Qy Query Match 100.0%; Score 10; DB 24; Length 22;
 Db Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 Db 7 CTTCTCTTTT 16

RESULT 47
 AAX10022/c
 ID AAX10022 standard; DNA; 23 BP.
 XX AC AAX10022;
 XX DT 24-MAR-1999 (first entry)
 XX DE Human biallelic polymorphic marker downstream primer #328.
 XX KW Polymorphism; biallelic; human; forensic; paternity testing; disease;
 KW detection; phenotypic typing; characteristic; infection; hereditary;
 KW autoimmune disease; cancer; inflammation; drug; therapy; medicament;
 XX treatment; marker; primer; ss.
 XX OS Synthetic.
 XX OS Homo sapiens.
 XX FN WO9820165-A2.
 XX PD 14-MAY-1998.
 XX PF 05-NOV-1997; 97WO-US20313.
 XX PR 06-NOV-1996; 96US-0030455.
 XX PA (WHEED) WHITEHEAD INST BIOMEDICAL RES.
 XX PI Hudson T, Lander ES, Wang D;
 XX WI; 1998-286974/25.
 XX New isolated nucleic acid segments from the human genome - used for
 PT determining polymorphic forms for use in e.g. forensics, paternity
 PT testing or phenotypic typing for disease
 XX Claim 16; Page 92; 310pp; English.

CC AAX09121-X10268 are allele-specific oligonucleotide primers used in the
 CC isolation of various biallelic polymorphic markers found in the human
 CC genome (represented in AAX10269-X12937). These primers can be used in a
 CC method for determining polymorphic forms in an individual for use in
 CC e.g. forensics, paternity testing or for phenotypic typing for diseases
 CC such as agammaglobulinemia, diabetes insipidus, Lesch-Nyhan syndrome,
 CC muscular dystrophy, Wiskott-Aldrich syndrome, Fabry's disease, familial
 CC hypercholesterolemia, polycystic kidney disease, hereditary
 CC spherocytosis, von Willebrand's disease, tuberous sclerosis, hereditary
 CC haemorrhagic telangiectasia, familial colonic polyposis, Ehlers-Danlos
 CC syndrome, osteogenesis imperfecta, acute intermittent porphyria,
 CC autoimmune diseases, inflammation, cancer, diseases of the nervous
 CC system, infection by pathogenic microorganisms, and characteristics such
 CC as longevity, appearance (e.g. baldness, obesity), strength, speed,
 CC endurance, fertility, and susceptibility or receptivity to particular
 CC drugs or therapeutic treatments. The isolated polymorphic nucleic acid
 CC segments can also be used to produce medicaments for the treatment or
 CC prophylaxis of such diseases.

```

SQ Sequence 23 BP; 16 A; 0 C; 7 G; 0 U; 0 other;
  Query Match      100.0%; Score 10; DB 19; Length 23;
  Best Local Similarity 100.0%; Pred. No. 1.8e+04;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

RESULT 48
AAQ50920/c
ID AAQ50920 standard; DNA; 24 BP.
XX
AC AAQ50920;
XX
DT 25-MAR-2003 (updated)
DT 19-MAY-1994 (first entry)
XX
DE T-cell antigen receptor V-beta14 PCR primer.
XX
XX RT-PCR; polymerase chain reaction; amplification; SSCP;
KW single-strand conformation polymorphism; variable domain;
KW subtype beta 14; ss.
XX
OS Synthetic.
XX
PN WO9322455-A1.
XX
PD 11-NOV-1993.
XX
PF 30-APR-1993; 93WO-JP00577.
XX
PR 30-APR-1992; 92JP-0111467.
PR 31-JUL-1992; 92JP-0205054.
XX
PA (LTLT-) LTT INST CO LTD.
PA (TAIS) TAISHO PHARM CO LTD.
XX
PI Ikeda Y, Mizushima Y, Nishioka K, Sakoda H, Yamamoto K;
DR WPI; 1993-368813/46.
XX
PT Detection of expression of T-cell antigen receptor gene - in
PT cancer, viral or immune disease patients, by polymerase chain
PT reaction amplification of the gene and SSCP analysis
XX
PS Claim 4; Page 13; 47pp; Japanese.
XX
CC Primers corresp. to DNA coding for part of the beta-chain of the T
CC cell antigen receptor (pref. the variable region primers AAQ50905-
CC AAQ50926) are used in PCR to amplify the T cell antigen receptor gene.
CC The amplified gene is detected by the single-strand conformation
CC polymorphism method using hybridisation probes corresp. to the
CC beta-chain J domain (see AAQ50928-Q50940).
CC (Updated on 25-MAR-2003 to correct FN field.)
XX
SQ Sequence 24 BP; 10 A; 3 C; 7 G; 4 T; 0 other;
  Query Match      100.0%; Score 10; DB 14; Length 24;
  Best Local Similarity 100.0%; Pred. No. 1.8e+04;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

RESULT 49
AAAT10390/c
ID AAAT10390 standard; cDNA; 24 BP.
XX

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AC AAT10390;
XX
DT 02-APR-1996 (first entry)
XX
DE T-cell receptor primer Vbeta14.
XX
KW T cell receptor; beta chain; variable region; rheumatoid arthritis;
KW synovial joint fluid; PCR; amplification; primer; immunogen; vaccine;
KW immune disease; ss.
XX
OS Synthetic.
XX
PN WO9523164-A1.
XX
PD 31-AUG-1995.
XX
PF 23-FEB-1995; 95WO-EP00670.
XX
PR 23-FEB-1994; 94EP-0200454.
XX
PA (ALKU ) AKZO NOBEL NV.
XX
PI Graus JPM, Rijnders AWM, Van Der Maaden JM;
XX WPI; 1995-311502/40.
XX
DR Peptide contained in the variable region of a T-cell receptor beta
PT chain - specifically associated with immune disease, esp. rheumatoid
PT arthritis
XX
PS Example 2; Page 24; 55pp; English.
XX
CC The primers AAT10352-97 were used to PCR amplify the T cell receptor
CC beta chain variable regions from T cell culture clones, isolated from
CC the synovial joint fluid of 11 patients suffering from rheumatoid
CC arthritis. The coding sequences were shown to contain the nucleotide
CC sequence AAT07409. The encoded polypeptide can be used as an immunogenic
CC cpd. for the detection of or predisposition to an immune disease, or for
CC use as a vaccine for prevention or treatment of an immune disease. This
CC primer amplifies the variable region from the 14.1 family of clones.
XX
SQ Sequence 24 BP; 10 A; 3 C; 7 G; 4 T; 0 other;
  Query Match      100.0%; Score 10; DB 16; Length 24;
  Best Local Similarity 100.0%; Pred. No. 1.8e+04;
  Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

RESULT 50
AAT98116/c
ID AAT98116 standard; DNA; 24 BP.
XX
AC AAT98116;
XX
DT 13-MAR-1998 (first entry)
XX
DE Primer V-beta(14) for T-cell receptor beta chain variable region.
XX
KW Antibody; T-cell receptor; beta chain; human immunodeficiency virus;
KW HIV; blood; attenuation; primer; PCR; amplification; variable region;
KW constant region; TCR; ss.
XX
OS Synthetic.
OS Homo sapiens.
XX
PN US5665355-A.
XX
PD 09-SEP-1997.
XX

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PF 07-JUN-1995; 95US-0488212.
 XX
 PR 09-NOV-1992; 92US-0973485.
 PR 18-OCT-1994; 94US-0408011.
 PR 07-JUN-1995; 95US-0488212.
 XX
 PA (CONS-) CONSORZIO BIOTECNOLOGIE.
 XX
 XX Primi D;
 XX WPI; 1997-456759/42.
 XX
 PT Removal of T-cell receptor-specific antibody from blood of
 PT HIV-infected person - by extracorporeal blood treatment, to
 PT attenuate or avert development of AIDS from HIV infection
 XX
 XX Example 1; Column 11; 43pp; English.
 XX
 CC The invention relates to a method for removing an antibody specific for
 CC TCR-V beta (T-cell receptor V beta protein) from an HIV-infected person
 CC by removing blood from the person, removing the antibody from the blood,
 CC and reintroducing the blood into the person, thus allowing attenuation
 CC or aversion of immunodeficiency. The primers AAT98100-T98150 are used
 CC to check the efficiency of removal by detecting expression of the
 CC TCR-V-beta and V-alpha genes in a blood sample after treatment. This
 CC primer is targeted to the variable region sequence and can be used in
 CC the amplification with primer AAT98100.
 XX
 SQ Sequence 24 BP; 10 A; 3 C; 7 G; 4 T; 0 other;
 Query Match 100.0%; Score 10; DB 18; Length 24;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 18 CTTCTCTTTT 9
 RESULT 51
 AAX85959/c
 ID AAX85959 standard; DNA; 24 BP.
 AC AAX85959;
 XX
 DT 13-SEP-1999 (first entry)
 XX
 DE PCR primer used to amplify T cell receptor beta-chain cDNA.
 XX
 KW Acquired immune deficiency syndrome; free antibody; paratope; epitope;
 KW T cell receptor variable beta region; TCR-V beta region; binding agent;
 KW CD4+ T cell; HIV; PCR primer; ss.
 XX
 OS Synthetic.
 OS Homo sapiens.
 XX
 PN US5928642-A.
 XX
 PD 27-JUL-1999.
 XX
 PF 18-OCT-1994; 94US-0408011.
 XX
 PR 09-NOV-1992; 92US-0973485.
 PR 18-OCT-1994; 94US-0408011.
 XX
 PA (CONS-) CONSORZIO BIOTECNOLOGIE.
 XX
 XX Primi D;
 XX WPI; 1999-429481/36.
 DR
 XX
 PT Diagnosis and treatment of acquired immune deficiency syndrome
 XX

PS Example 1; Column 11; 42pp; English.

XX The specification describes a method for the diagnosis and treatment of
 CC acquired immune deficiency syndrome, in a person having free antibodies
 CC which have a paratope capable of binding to an epitope of a T cell
 CC receptor variable beta (TCR-V beta) region. The method comprises
 CC administering a binding agent homologous with the TCR-V beta
 CC epitope. The binding agent is useful in assays for detecting various
 CC CD4+ T cell subpopulations which carry particular V beta components.
 CC The binding agent is also useful in the treatment of people infected
 CC with HIV where it is able to remove an antibody able to bind with an
 CC epitope on a TCR-V beta cell in the blood of an infected person. PCR
 CC primers AAX85943-71 represent T cell receptor beta-chain primers used
 CC in the course of the invention.

SQ Sequence 24 BP; 10 A; 3 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 20; Length 24;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 Db 18 CTTCTCTTTT 9

RESULT 52

AAX88135/c
 ID AAX88135 standard; DNA; 24 BP.

AC AAX88135;

XX
 DT 09-SEP-1999 (first entry)

XX T cell receptor beta chain primer V-beta14.

XX T cell receptor; beta chain; primer; antibody; paratope; AIDS; vaccine;
 KW epitope; TCR-V beta; immunogenic; anti-idiotypic; antiviral; detection;
 KW CD4+ cell subpopulation; acquired immune deficiency syndrome; ss.

OS Synthetic.

XX US5925513-A.

XX 20-JUL-1999.

XX 07-JUN-1995; 95US-0488209.

XX 09-NOV-1992; 92US-0973485.

PR 18-OCT-1994; 94US-0408011.

PR 07-JUN-1995; 95US-0488209.

XX (CONS-) CONSORZIO BIOTECNOLOGIE.

XX Primi D;

XX WPI; 1999-418267/35.

XX Diagnosis and treatment of acquired immune deficiency syndrome onset

PS Example 1; Column 11-12; 42pp; English.

XX This invention describes novel method for binding free antibodies having
 CC a paratope specific to an epitope on a T cell receptor (TCR-V beta)
 CC while providing an immunogenic substance able to raise anti-idiotypic
 CC antibodies which bind to free antibodies bound at the same paratope
 CC specific to the epitope on the TCR-V beta and introducing this into a
 CC person to raise anti-idiotypic antibodies. The products of the invention
 CC have antiviral activity and can be used in vaccines. The specific
 CC antibody binding affinities are useful in assays which detect the
 CC presence of CD4+ cell subpopulations carrying particular V beta
 CC components of the TCR-V beta in people infected with acquired immune
 CC deficiency syndrome (AIDS). AAX88119-X88169 represents primers used in

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CC the method of the invention.
XX
SQ Sequence 24 BP; 10 A; 3 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 20; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 18 CTTCTCTTTT 9

RESULT 53
AAI99892/c
ID AAI99892 standard; DNA; 24 BP.
XX
AC AAI99892;
XX
DT 30-JAN-2002 (first entry)
XX
DE Human dihydroorotase 11 PCR primer SEQ ID NO 4.
XX
KW Human; dihydroorotase 11; cytostatic; virucidal; immunomodulatory;
KW antiinflammatory; haemostatic; malignant tumour; HIV; infection;
KW human immunodeficiency virus; immunological disease; gene therapy;
KW PCR primer; ss.
XX
OS Homo sapiens.
XX
PN WO200173054-A1.
XX
PD 04-OCT-2001.
XX
PF 26-MAR-2001; 2001WO-CN00421.
XX
PR 27-MAR-2000; 2000CN-0115157.
XX
PA (SHAN-) SHANGHAI BIOWINDOW GENE DEV INC.
XX
PI Mao Y, Xie Y;
XX
DR WPI; 2001-602867/68.
XX
PT New human dihydroorotase 11 for diagnosing and treating malignant
PT tumour, haemopathy, human immunodeficiency virus infection, immunological
PT diseases and various inflammations -
XX
PS Example 2; Page 17; 34pp; Chinese.
XX
CC The invention relates to human dihydroorotase 11 with cytostatic,
CC virucidal, immunomodulatory, antiinflammatory and haemostatic
CC activity. The protein and encoding polynucleotide are used in diagnosis
CC and treatment of malignant tumour, haemopathy, human immunodeficiency
CC virus (HIV) infection, immunological diseases and various
CC inflammations. The polynucleotide is useful in gene therapy. The present
CC sequence is that of a PCR primer, useful to the invention.
XX
SQ Sequence 24 BP; 8 A; 3 C; 8 G; 5 T; 0 other;

Query Match 100.0%; Score 10; DB 22; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 10 CTTCTCTTTT 1

RESULT 54
AAH19640/c
ID AAI99640 standard; DNA; 24 BP.
XX

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AC AAI9640;
XX
DT 01-AUG-2001 (first entry)
XX
DE Melon fusarium wilt-resistance marker FM forward primer FM-1.
XX
KW Melon; cucurbit; Fusarium wilt; infection; resistance; susceptibility;
KW genotype identification; polymerase chain reaction; PCR;
KW Fusarium oxysporum; marker; PCR primer; ss.
XX
OS Cucumis melo.
XX
PN WO200127332-A1.
XX
PD 19-APR-2001.
XX
PF 13-OCT-2000; 2000WO-US28633.
XX
PR 13-OCT-1999; 99US-0417722.
XX
PA (UYCL-) UNIV CLEMSON.
XX
PI Dean RA, Wang Y;
XX
DR WPI; 2001-300220/31.
XX
PT Identifying cucurbit genotypes to determine susceptibility or
PT resistance to Fusarium wilt disease, comprises using the polymerase
PT chain reaction (PCR) and comparing the PCR product with known
PT resistant/susceptible genotypes -
XX
PS Claim 10; Fig 2; 36pp; English.
XX
CC The present sequence is a primer used in a method for identifying
CC genotypes in cucurbit plants, particularly species of melon, that
CC are resistant or susceptible to Fusarium wilt disease. The method
CC comprises extracting genomic DNA from a plant sample and using
CC polymerase chain reaction (PCR) to produce DNA products for
CC comparison with known DNA sequences of Fusarium-resistant/susceptible
CC genotypes. The method is especially useful for establishing whether
CC cucurbit plants are resistant or susceptible to Fusarium wilt
CC disease caused by Fusarium oxysporum infection. The method uses PCR
CC to detect the genotypes. PCR is sensitive and accurate, and it is a
CC much more rapid procedure than previous methods such as artificial
CC inoculation (AI). The new method is also more cost effective and
CC more suitable for high throughput analysis.
CC Note: The present sequence is given as SEQ ID NO: 3 in Figure 2,
CC but it is different to the sequence given as SEQ ID NO: 3 in the
CC sequence listing.
XX
SQ Sequence 24 BP; 14 A; 1 C; 8 G; 1 T; 0 other;

Query Match 100.0%; Score 10; DB 22; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 23 CTTCTCTTTT 14

RESULT 55
ABS71673/c
ID ABS71673 standard; DNA; 24 BP.
XX
AC ABS71673;
XX
DT 28-NOV-2002 (first entry)
XX
DE T cell receptor (TCR) variable beta (BV) peptide RT-PCR primer #16.
XX
KW T cell receptor; TCR; receptor; variable beta peptide; BV peptide; TCRV;
KW T cell variable gene; T cell regulatory activity; autoimmune disease;

```

KW multiple sclerosis; human; reverse transcriptase; RT-PCR; primer; ss.
 XX Homo sapiens.
 OS US2002107388-A1.
 PN 08-AUG-2002.
 PD 10-MAY-2001; 2001US-0853830.
 PP 12-MAY-2000; 2000US-203984P.
 PR (VAND/) VANDENBARK A A.
 XX Vandenbark AA;
 PI WPI; 2002-697882/75.
 DR Identifying a T cell receptor variable gene expressed by target T cells
 XX in an individual is useful to identify disease-associated T cells for
 PT design of individualised therapies, particularly for autoimmune disease
 PT -
 PS Example 2; Page 11; 20pp; English.
 XX The invention relates to a method for identifying a T cell receptor
 CC variable (TCRV) gene expressed by target T cells in an individual,
 CC comprising determining expression of TCRV genes by activated T cells from
 CC the individual and determining regulatory activity elicited in response
 CC to TCRV peptides from the individual. A preferentially expressed TCRV
 CC gene whose TCRV peptide elicits low T cell regulatory activity is
 CC identified as a variable gene expressed by target T cells. The method is
 CC used to identify disease-associated T cells in an individual so that
 CC individualised therapies can be designed to prevent or treat the disease,
 CC particularly an autoimmune disease, especially multiple sclerosis. This
 CC sequence represents a reverse transcriptase PCR (RT-PCR) primer used in
 CC analysis of expression of DNA encoding TCR variable beta (BV) peptides.
 XX Sequence 24 BP; 10 A; 3 C; 7 G; 4 T; 0 other;
 SQ Query Match 100.0%; Score 10; DB 24; Length 24;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 18 CTTCTCTTTT 9
 RESULT 56
 ABK67667/C
 ID ABK67667 standard; DNA; 24 BP.
 XX AC ABK67667;
 AC 02-JUL-2002 (first entry)
 DT B. subtilis knock-out PCR primer KO murA R1.
 DE Antibacterial; antiprotozoal; antifungal; cell growth inhibitor;
 XX KW protozoa; bacterium; fungus; amoeba; mycoplasma; murA;
 KW folA; yibD; primer; ss.
 XX OS Bacillus subtilis.
 OS WO200216940-A2.
 PN 28-FEB-2002.
 PD 23-AUG-2001; 2001WO-US26322.
 PP 23-AUG-2000; 2000US-226896P.
 PR

PA (GENO-) GENOME THERAPEUTICS CORP.
 PI Sulavik M, Ling LL, Opperman T, Moir DT, Bunker C;
 XX WPI; 2002-329705/36.
 DR Identification of a molecular target of a cell growth inhibitor,
 PT comprises target prediction processes to identify at least one
 PT modulated gene or gene product -
 XX Example 2; Page 97; 157pp; English.
 PS The invention relates to a method of identifying a molecular target of a
 XX cell growth inhibiting compound by: (a) identifying a compound or
 CC composition inhibiting growth in a population of cells; (b) performing
 CC target prediction processes to identify a modulated gene or gene product;
 CC and (c) comparing by target prediction processes with the gene or gene
 CC product identified by other prediction processes. The method is used for
 CC identifying the molecular target of a cell growth inhibitor. The cells
 CC are selected from organisms such as protozoa, bacterium, fungus, amoeba
 CC and mycoplasma. The method reaps the benefits of both cell-based and
 CC target-based screening. The method relies first on identification of
 CC compounds with good whole cell activity and then provides methods to
 CC identify the target of the compounds. Each of the methods is successful
 CC in isolation, but in combination of at least two (preferably at least
 CC three) provides a higher success in identifying the molecular target.
 CC ABK67622-ABK67679 represent PCR primers used in examples which
 CC demonstrate the method of the invention.
 XX Sequence 24 BP; 9 A; 4 C; 6 G; 5 T; 0 other;
 SQ Query Match 100.0%; Score 10; DB 24; Length 24;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 16 CTTCTCTTTT 7
 RESULT 57
 ABA04339/C
 ID ABA04339 standard; DNA; 24 BP.
 XX AC ABA04339;
 AC 05-MAR-2002 (first entry)
 DT Human BOLA structural domain zinc finger protein 37 PCR primer 1.
 DE Human; BOLA structural domain zinc finger protein 37; malignant tumour;
 XX KW nosohaemia; HIV infection; immunological disease; inflammation;
 KW PCR primer; ss.
 XX OS Homo sapiens.
 OS CN1307053-A.
 PN 08-AUG-2001.
 PD 28-JAN-2000; 2000CN-0111611.
 PP 28-JAN-2000; 2000CN-0111611.
 PR (BODA-) BODAO GENE TECH CO LTD SHANGHAI.
 PA Mao Y, Xie Y;
 XX WPI; 2002-062745/09.
 DR Polypeptide-human BOLA structural domain zinc finger protein 37 and
 PT polynucleotide for said polypeptide -
 XX

PS Example 3; Page 17 (Disclosure); 35pp; Chinese.

XX The present invention describes human BOLA structural domain zinc finger
 CC protein 37. (i). (I) can be used in the treatment of various diseases,
 CC such as malignant tumour, nosohaemia, HIV infection, immunological
 CC diseases and inflammations. The present sequence represents a PCR primer
 CC for human BOLA structural domain zinc finger protein 37, which is used
 CC in an example from the present invention.

XX Sequence 24 BP; 12 A; 4 C; 4 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 24;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 Db 18 CTTCTCTTTT 9

RESULT 59

ABL45589/C

ID ABL45589 standard; DNA; 24 BP.

XX AC ABL45589;

XX 11-APR-2002 (first entry)

DE Human chromosome 21q22.1 PCR primer SEQ ID NO:2633.

KW Human; chromosome 1p36-35; chromosome 21q22.1; genetic analysis;
 KW genome; PCR primer; ss.

OS Homo sapiens.

XX JP2001321190-A.

XX 20-NOV-2001.

XX 12-MAR-2001; 2001JP-0068285.

XX 10-MAR-2000; 2000JP-0066716.

PA (RIKA) RIKAGAKU KENKYUSHO.

XX (GENO-) GENOTEX YG.

XX WPI; 2002-144136/19.

XX Arraying genome clones -

PS Claim 6; Page 57; 528pp; Japanese.

XX The present invention describes a method of arraying genome clones. The
 CC method comprises: (a) clones of the genomic libraries contained in
 CC multiwell plates numbered for discrimination are mixed in each of the
 CC multiwell plates; (b) a primer designed based on the chromosome marker
 CC sequence is added to the mixture to carry out an amplification reaction;
 CC (c) a signal corresponding to the marker is detected from the resultant
 CC amplified product to specify the discrimination Nos. of the multiwell
 CC plates containing the clones having said marker sequence; (d) the order
 CC of the markers is changed so that the same discrimination Nos. succeed to
 CC the maximum in the specified discrimination Nos. to array the multiwell
 CC plates; (e) the clones in the specified discrimination Nos. to array the multiwell
 CC discrimination Nos. are mixed respectively in each wells of longitudinal
 CC and lateral directions; (f) the mixed clones are cultured and the
 CC resultant cultures are amplified by using the above primer; (g) signals
 CC are detected from the amplified products; (h) the clones in the multiwell
 CC plates are specified from the detected result; and (i) the clones are
 CC reconstituted as the positions on the chromosome and arrayed. The
 CC microarray is useful for gene analysis. ABL42957 to ABL45322 represent
 CC PCR primers for human chromosome 1p36-35 DNA, and ABL45323 to ABL45634
 CC represent PCR primers for human chromosome 21q22.1, which are
 CC specifically claimed for use in the present invention.

XX Sequence 24 BP; 12 A; 2 C; 6 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 24;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 Db 19 CTTCTCTTTT 10

RESULT 59

AAT76780

ID AAT76780 standard; DNA; 25 BP.

XX AC AAT76780;

XX 15-SEP-1997 (first entry)

XX Staphylococcus aureus exfoliative toxin A competitor primer ETA-B2.

XX Asymmetric polymerase chain reaction; nucleic acid amplification;
 KW PCR; detection; assay; exfoliative toxin A; ETA; skin lesion;
 KW competitive primer; capture probe; ss.

XX Synthetic.

XX US5627054-A.

XX 06-MAY-1997.

XX 05-APR-1996; 96US-0628417.

XX 05-APR-1996; 96US-0628417.

XX (USSA) US SEC OF ARMY.

XX Gillespie D;

XX WPI; 1997-271311/24.

XX Quantitative nucleic acid amplification - by competitor primer
 PT asymmetric polymerase chain reaction

XX Example 1; Column 5; 9pp; English.

XX In a specific example of a novel process for amplifying an amount
 CC (known or unknown) of a double-stranded nucleic acid segment to produce
 CC single-stranded nucleic acid in an amount that is proportional to the
 CC starting amount of the nucleic acid, the Staphylococcus aureus
 CC exfoliative toxin A (ETA) gene was used as the DNA template. The
 CC region comprising nucleotides 165-436 was amplified by symmetric,
 CC asymmetric or competitor primer asymmetric PCR using the primers
 CC ETA-A2 and ETA-B (see AAT76778 and AAT76779). For asymmetric PCR, the
 CC amount of primer ETA-B was reduced and for competitor primer
 CC asymmetric PCR a competitor primer ETA-B2 (see AAT76780) was added
 CC with upstream primer ETA-A2 after the initial cycling reaction. PCR
 CC products containing ETA-specific sequences were detected
 CC radioactively by a capture system which employed a bifunctional
 CC capture probe ETA-CP (see AAT76781 and AAT76782). ETA-CP was designed
 CC to capture the amplified sense strand onto capture membranes
 CC through hybridisation between the first 40 nucleotides of ETA-CP
 CC and nucleotides 321-360 of the ETA gene and through hybridisation
 CC of the poly(dA) tail on ETA-CP with poly(dT) tails on the capture
 CC membranes. A radioactively labelled "label probe" (see AAT76783),
 CC complementary to nucleotides 389-410 of the ETA gene was used to
 CC detect the amplicons. Results showed that hybridisation of the
 CC capture probe and label probe to the denatured symmetric PCR
 CC product was much less efficient than hybridisation to the
 CC single-stranded PCR products of the asymmetric and competitor
 CC primer asymmetric reactions.

XX

```
SQ Sequence 25 BP; 4 A; 5 C; 3 G; 8 T; 5 other;
Query Match 100.0%; Score 10; DB 18; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17

RESULT 60
ABV82471/c
ID ABV82471 standard; DNA; 25 BP.
XX AC ABV82471;
XX DT 03-JAN-2003 (first entry)
XX DE Human HTPL scanning oligonucleotide SEQ ID 3717.
XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
KW human testis expressed Patched like protein; testis; adrenal; liver;
KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
XX OS Homo sapiens.
XX PN EP1229046-A2.
XX PD 07-AUG-2002.
XX PF 28-JAN-2002; 2002EP-0001167.
XX PR 30-JAN-2001; 2001WO-US00663.
XX PR 30-JAN-2001; 2001WO-US00664.
XX PR 30-JAN-2001; 2001WO-US00665.
XX PR 30-JAN-2001; 2001WO-US00666.
XX PR 30-JAN-2001; 2001WO-US00667.
XX PR 30-JAN-2001; 2001WO-US00668.
XX PR 23-MAY-2001; 2001US-0864761.
XX PR 09-OCT-2001; 2001US-0327898.
XX PA (AEOM-) AEOMICA INC.
XX PI Zhan J;
XX DR WPI; 2002-676582/73.
XX PT Novel isolated human testis expressed Patched like protein (HTPL),
XX PT useful for identifying agonist and antagonist and specific binding
XX PT partners, and for treating subjects having defects in HTPL -
XX PS Example 2; Page 551; 718pp; English.
XX CC The present invention relates to human testis expressed Patched like
XX CC protein (HTPL, see ABV78759 to ABV78762 and ABB98520). HTPL
XX CC has two isoforms, with a few single base pair differences between the
XX CC two. One of the single base pair changes introduces a premature stop
XX CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
XX CC shares an overall structure organisation with the Patched protein. The
XX CC shared structural features strongly imply that HTPL plays a role similar
XX CC to that of Patched, and is a potential tumour suppressor. HTPL is
XX CC important in regulating male germ cell development, and the HTPL gene was
XX CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
XX CC useful for diagnosing a disorder caused by mutation in HTPL, and in
XX CC therapy and manufacture of a medicament for treatment or prevention of
XX CC such disorder associated with decreased expression or activity of human
XX CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
XX CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
XX CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
XX CC clinically useful diagnostic markers and potential therapeutic agents for
XX CC male infertility and cancer. The present oligonucleotide was used in an
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CC example from the invention.
SQ Sequence 25 BP; 9 A; 2 C; 9 G; 5 T; 0 other;
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16

RESULT 61
ABV82472/c
ID ABV82472 standard; DNA; 25 BP.
XX AC ABV82472;
XX DT 03-JAN-2003 (first entry)
XX DE Human HTPL scanning oligonucleotide SEQ ID 3718.
XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
KW human testis expressed Patched like protein; testis; adrenal; liver;
KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
XX OS Homo sapiens.
XX PN EP1229046-A2.
XX PD 07-AUG-2002.
XX PF 28-JAN-2002; 2002EP-0001167.
XX PR 30-JAN-2001; 2001WO-US00663.
XX PR 30-JAN-2001; 2001WO-US00664.
XX PR 30-JAN-2001; 2001WO-US00665.
XX PR 30-JAN-2001; 2001WO-US00666.
XX PR 30-JAN-2001; 2001WO-US00667.
XX PR 30-JAN-2001; 2001WO-US00668.
XX PR 23-MAY-2001; 2001US-0864761.
XX PR 09-OCT-2001; 2001US-0327898.
XX PA (AEOM-) AEOMICA INC.
XX PI Zhan J;
XX DR WPI; 2002-676582/73.
XX PT Novel isolated human testis expressed Patched like protein (HTPL),
XX PT useful for identifying agonist and antagonist and specific binding
XX PT partners, and for treating subjects having defects in HTPL -
XX PS Example 2; Page 551; 718pp; English.
XX CC The present invention relates to human testis expressed Patched like
XX CC protein (HTPL, see ABV78759 to ABV78762 and ABB98520). HTPL
XX CC has two isoforms, with a few single base pair differences between the
XX CC two. One of the single base pair changes introduces a premature stop
XX CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
XX CC shares an overall structure organisation with the Patched protein. The
XX CC shared structural features strongly imply that HTPL plays a role similar
XX CC to that of Patched, and is a potential tumour suppressor. HTPL is
XX CC important in regulating male germ cell development, and the HTPL gene was
XX CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
XX CC useful for diagnosing a disorder caused by mutation in HTPL, and in
XX CC therapy and manufacture of a medicament for treatment or prevention of
XX CC such disorder associated with decreased expression or activity of human
XX CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
XX CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
XX CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
XX CC clinically useful diagnostic markers and potential therapeutic agents for
XX CC male infertility and cancer. The present oligonucleotide was used in an
```

CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.
 SQ Sequence 25 BP; 8 A; 2 C; 9 G; 6 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 CTTCTCTTTT 10
 |||||
 Db 24 CTTCTCTTTT 15

RESULT 62
 ABV82473/c
 ID ABV82473 standard; DNA; 25 BP.
 XX
 AC ABV82473;
 XX
 DT 03-JAN-2003 (first entry)
 XX
 DE Human HTPL scanning oligonucleotide SEQ ID 3719.
 XX
 DE Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
 OS Homo sapiens.
 XX
 XX Homo sapiens.
 PN EP1229046-A2.
 XX
 XX EP1229046-A2.
 PD 07-AUG-2002.
 XX
 XX 28-JAN-2002; 2002EP-0001167.
 PF
 PR 30-JAN-2001; 2001WO-US00663.
 PR 30-JAN-2001; 2001WO-US00664.
 PR 30-JAN-2001; 2001WO-US00665.
 PR 30-JAN-2001; 2001WO-US00666.
 PR 30-JAN-2001; 2001WO-US00667.
 PR 30-JAN-2001; 2001WO-US00668.
 PR 23-MAY-2001; 2001US-0864761.
 PR 09-OCT-2001; 2001US-0327898.
 XX
 PA (AEOM-) AEOMICA INC.
 XX
 PI Zhan J;
 XX
 DR WPI; 2002-676582/73.
 XX
 PT Novel isolated human testis expressed Patched like protein (HTPL),
 PT useful for identifying agonist and antagonist and specific binding
 PT partners, and for treating subjects having defects in HTPL -
 XX
 PS Example 2; Page 551; 718pp; English.
 XX
 CC The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
 CC shares an overall structure organisation with the Patched protein. The
 CC shared structural features strongly imply that HTPL plays a role similar
 CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC therapy and manufacture of a medicament for treatment or prevention of
 CC such disorder associated with decreased expression or activity of human
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and

CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.
 SQ Sequence 25 BP; 9 A; 2 C; 9 G; 5 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 CTTCTCTTTT 10
 |||||
 Db 23 CTTCTCTTTT 14

RESULT 63
 ABV82474/c
 ID ABV82474 standard; DNA; 25 BP.
 XX
 AC ABV82474;
 XX
 DT 03-JAN-2003 (first entry)
 XX
 DE Human HTPL scanning oligonucleotide SEQ ID 3720.
 XX
 DE Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
 OS Homo sapiens.
 XX
 XX Homo sapiens.
 PN EP1229046-A2.
 XX
 XX EP1229046-A2.
 PD 07-AUG-2002.
 XX
 XX 28-JAN-2002; 2002EP-0001167.
 PF
 PR 30-JAN-2001; 2001WO-US00663.
 PR 30-JAN-2001; 2001WO-US00664.
 PR 30-JAN-2001; 2001WO-US00665.
 PR 30-JAN-2001; 2001WO-US00666.
 PR 30-JAN-2001; 2001WO-US00667.
 PR 30-JAN-2001; 2001WO-US00668.
 PR 23-MAY-2001; 2001US-0864761.
 PR 09-OCT-2001; 2001US-0327898.
 XX
 PA (AEOM-) AEOMICA INC.
 XX
 PI Zhan J;
 XX
 DR WPI; 2002-676582/73.
 XX
 PT Novel isolated human testis expressed Patched like protein (HTPL),
 PT useful for identifying agonist and antagonist and specific binding
 PT partners, and for treating subjects having defects in HTPL -
 XX
 PS Example 2; Page 551; 718pp; English.
 XX
 CC The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
 CC shares an overall structure organisation with the Patched protein. The
 CC shared structural features strongly imply that HTPL plays a role similar
 CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC therapy and manufacture of a medicament for treatment or prevention of
 CC such disorder associated with decreased expression or activity of human
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and

CC such disorder associated with decreased expression or activity of human
CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
CC clinically useful diagnostic markers and potential therapeutic agents for
CC male infertility and cancer. The present oligonucleotide was used in an
CC example from the invention.
XX
SQ Sequence 25 BP; 9 A; 2 C; 9 G; 5 T; 0 other;
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 22 CTTCTCTTTT 13
RESULT 64
ABV82475/C
ID ABV82475 standard; DNA; 25 BP.
XX AC ABV82475;
XX XX
DT 03-JAN-2003 (first entry)
XX
DE Human HTPL scanning oligonucleotide SEQ ID 3721.
XX
KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
KW human testis expressed Patched like protein; testis; adrenal; liver;
KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
XX
OS Homo sapiens.
XX
PN EP1229046-A2.
XX
PD 07-AUG-2002.
XX
PF 28-JAN-2002; 2002EP-0001167.
XX
PR 30-JAN-2001; 2001WO-US000663.
PR 30-JAN-2001; 2001WO-US000664.
PR 30-JAN-2001; 2001WO-US000665.
PR 30-JAN-2001; 2001WO-US000667.
PR 30-JAN-2001; 2001WO-US000668.
PR 30-JAN-2001; 2001WO-US000669.
PR 23-MAY-2001; 2001US-0864761.
PR 09-OCT-2001; 2001US-0327898.
XX
PA (AEOM-) AEOMICA INC.
XX
PI Zhan J;
XX
DR WPI; 2002-676582/73.
XX
PT Novel isolated human testis expressed Patched like protein (HTPL),
PT useful for identifying agonist and antagonist and specific binding
PT partners, and for treating subjects having defects in HTPL -
XX
PS Example 2; Page 551; 718pp; English.
XX
CC The present invention relates to human testis expressed Patched like
CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
CC has two isoforms, with a few single base pair differences between the
CC two. One of the single base pair changes introduces a premature stop
CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
CC shares an overall structure organisation with the Patched protein. The
CC shared structural features strongly imply that HTPL plays a role similar
CC to that of Patched, and is a potential tumour suppressor. HTPL is
CC important in regulating male germ cell development, and the HTPL gene was
CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are

CC useful for diagnosing a disorder caused by mutation in HTPL, and in
CC therapy and manufacture of a medicament for treatment or prevention of
CC such disorder associated with decreased expression or activity of human
CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
CC clinically useful diagnostic markers and potential therapeutic agents for
CC male infertility and cancer. The present oligonucleotide was used in an
CC example from the invention.
XX
SQ Sequence 25 BP; 9 A; 2 C; 8 G; 6 T; 0 other;
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 21 CTTCTCTTTT 12
RESULT 65
ABV82476/C
ID ABV82476 standard; DNA; 25 BP.
XX AC ABV82476;
XX XX
DT 03-JAN-2003 (first entry)
XX
DE Human HTPL scanning oligonucleotide SEQ ID 3722.
XX
KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
KW human testis expressed Patched like protein; testis; adrenal; liver;
KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
XX
OS Homo sapiens.
XX
PN EP1229046-A2.
XX
PD 07-AUG-2002.
XX
PF 28-JAN-2002; 2002EP-0001167.
XX
PR 30-JAN-2001; 2001WO-US000663.
PR 30-JAN-2001; 2001WO-US000664.
PR 30-JAN-2001; 2001WO-US000665.
PR 30-JAN-2001; 2001WO-US000667.
PR 30-JAN-2001; 2001WO-US000668.
PR 30-JAN-2001; 2001WO-US000669.
PR 23-MAY-2001; 2001US-0864761.
PR 09-OCT-2001; 2001US-0327898.
XX
PA (AEOM-) AEOMICA INC.
XX
PI Zhan J;
XX
DR WPI; 2002-676582/73.
XX
PT Novel isolated human testis expressed Patched like protein (HTPL),
PT useful for identifying agonist and antagonist and specific binding
PT partners, and for treating subjects having defects in HTPL -
XX
PS Example 2; Page 551; 718pp; English.
XX
CC The present invention relates to human testis expressed Patched like
CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
CC has two isoforms, with a few single base pair differences between the
CC two. One of the single base pair changes introduces a premature stop
CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
CC shares an overall structure organisation with the Patched protein. The
CC shared structural features strongly imply that HTPL plays a role similar
CC to that of Patched, and is a potential tumour suppressor. HTPL is
CC important in regulating male germ cell development, and the HTPL gene was
CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are

CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC therapy and manufacture of a medicament for treatment or prevention of
 CC such disorder associated with decreased expression or activity of human
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.
 XX
 SQ Sequence 25 BP; 9 A; 2 C; 8 G; 6 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTTTT 10
 |||||
 Db 20 CTCTCTTTT 11
 |||||

RESULT 66
 ABV82477/c
 ID ABV82477 standard; DNA; 25 BP.

XX AC ABV82477;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 3723.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EP1229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US006663.

XX PR 30-JAN-2001; 2001WO-US006664.

XX PR 30-JAN-2001; 2001WO-US006665.

XX PR 30-JAN-2001; 2001WO-US006667.

XX PR 30-JAN-2001; 2001WO-US006668.

XX PR 23-MAY-2001; 2001US-0864761.

XX PR 09-OCT-2001; 2001US-0327898.

XX PA (AEOM-) AEOMICA INC.

XX PI Zhan J;

XX DR WPI; 2002-676582/73.

XX PT Novel isolated human testis expressed Patched like protein (HTPL),
 PT useful for identifying agonist and antagonist and specific binding
 PT partners, and for treating subjects having defects in HTPL -
 XX
 XX Example 2; Page 552; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
 CC shares an overall structure organisation with the Patched protein. The

CC shared structural features strongly imply that HTPL plays a role similar
 CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC therapy and manufacture of a medicament for treatment or prevention of
 CC such disorder associated with decreased expression or activity of human
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.
 XX

SQ Sequence 25 BP; 9 A; 1 C; 8 G; 7 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTTTT 10
 |||||
 Db 19 CTCTCTTTT 10
 |||||

RESULT 67
 ABV82478/c
 ID ABV82478 standard; DNA; 25 BP.

XX AC ABV82478;

XX DT 03-JAN-2003 (first entry)

XX DE Human HTPL scanning oligonucleotide SEQ ID 3724.

XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX OS Homo sapiens.

XX PN EP1229046-A2.

XX PD 07-AUG-2002.

XX PF 28-JAN-2002; 2002EP-0001167.

XX PR 30-JAN-2001; 2001WO-US006663.

XX PR 30-JAN-2001; 2001WO-US006664.

XX PR 30-JAN-2001; 2001WO-US006665.

XX PR 30-JAN-2001; 2001WO-US006667.

XX PR 30-JAN-2001; 2001WO-US006668.

XX PR 23-MAY-2001; 2001US-0864761.

XX PR 09-OCT-2001; 2001US-0327898.

XX PA (AEOM-) AEOMICA INC.

XX PI Zhan J;

XX DR WPI; 2002-676582/73.

XX PT Novel isolated human testis expressed Patched like protein (HTPL),
 PT useful for identifying agonist and antagonist and specific binding
 PT partners, and for treating subjects having defects in HTPL -
 XX
 XX Example 2; Page 552; 718pp; English.

XX CC The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop

CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
CC shares an overall structure organisation with the Patched protein. The
CC shared structural features strongly imply that HTPL plays a role similar
CC to that of Patched, and is a potential tumour suppressor. HTPL is
CC important in regulating male germ cell development, and the HTPL gene was
CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
CC useful for diagnosing a disorder caused by mutation in HTPL, and in
CC therapy and manufacture of a medicament for treatment or prevention of
CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
CC clinically useful diagnostic markers and potential therapeutic agents for
CC male infertility and cancer. The present oligonucleotide was used in an
CC example from the invention.

XX SQ Sequence 25 BP; 9 A; 1 C; 9 G; 6 T; 0 other;
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

RESULT 68
ABV82479/c
ID ABV82479 standard; DNA; 25 BP.
XX AC ABV82479;
XX DT 03-JAN-2003 (first entry)
XX DE Human HTPL scanning oligonucleotide SEQ ID 3725.
XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
XX KW human testis expressed Patched like protein; testis; adrenal; liver;
XX KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
XX KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
XX OS Homo sapiens.
XX PN EP1229046-A2.
XX PD 07-AUG-2002.
XX PF 28-JAN-2002; 2002EP-0001167.
XX PR 30-JAN-2001; 2001WO-US00663.
XX PR 30-JAN-2001; 2001WO-US00664.
XX PR 30-JAN-2001; 2001WO-US00665.
XX PR 30-JAN-2001; 2001WO-US00667.
XX PR 30-JAN-2001; 2001WO-US00668.
XX PR 30-JAN-2001; 2001WO-US00669.
XX PR 23-MAY-2001; 2001US-084761.
XX PR 09-OCT-2001; 2001US-0327898.
XX PA (AEOM-) AEOMICA INC.
XX PI Zhan J;
XX WPI; 2002-676582/73.
XX DR Novel isolated human testis expressed Patched like protein (HTPL),
XX PT useful for identifying agonist and antagonist and specific binding
XX PT partners, and for treating subjects having defects in HTPL -
XX PS Example 2; Page 552; 718pp; English.
XX CC The present invention relates to human testis expressed Patched like
XX protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL

CC has two isoforms, with a few single base pair differences between the
CC two. One of the single base pair changes introduces a premature stop
CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
CC shares an overall structure organisation with the Patched protein. The
CC shared structural features strongly imply that HTPL plays a role similar
CC to that of Patched, and is a potential tumour suppressor. HTPL is
CC important in regulating male germ cell development, and the HTPL gene was
CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
CC useful for diagnosing a disorder caused by mutation in HTPL, and in
CC therapy and manufacture of a medicament for treatment or prevention of
CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
CC clinically useful diagnostic markers and potential therapeutic agents for
CC male infertility and cancer. The present oligonucleotide was used in an
CC example from the invention.

XX SQ Sequence 25 BP; 9 A; 2 C; 9 G; 5 T; 0 other;
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 17 CTTCTCTTTT 8

RESULT 69
ABV82480/c
ID ABV82480 standard; DNA; 25 BP.
XX AC ABV82480;
XX DT 03-JAN-2003 (first entry)
XX DE Human HTPL scanning oligonucleotide SEQ ID 3726.
XX KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
XX KW human testis expressed Patched like protein; testis; adrenal; liver;
XX KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
XX KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
XX OS Homo sapiens.
XX PN EP1229046-A2.
XX PD 07-AUG-2002.
XX PF 28-JAN-2002; 2002EP-0001167.
XX PR 30-JAN-2001; 2001WO-US00663.
XX PR 30-JAN-2001; 2001WO-US00664.
XX PR 30-JAN-2001; 2001WO-US00665.
XX PR 30-JAN-2001; 2001WO-US00667.
XX PR 30-JAN-2001; 2001WO-US00668.
XX PR 30-JAN-2001; 2001WO-US00669.
XX PR 23-MAY-2001; 2001US-084761.
XX PR 09-OCT-2001; 2001US-0327898.
XX PA (AEOM-) AEOMICA INC.
XX PI Zhan J;
XX WPI; 2002-676582/73.
XX DR Novel isolated human testis expressed Patched like protein (HTPL),
XX PT useful for identifying agonist and antagonist and specific binding
XX PT partners, and for treating subjects having defects in HTPL -
XX PS Example 2; Page 552; 718pp; English.
XX CC The present invention relates to human testis expressed Patched like
XX protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL

CC The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
 CC shares an overall structure organisation with the Patched protein. The
 CC shared structural features strongly imply that HTPL plays a role similar
 CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC such disorder associated with decreased expression or activity of human
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.

XX
 SQ Sequence 25 BP; 9 A; 3 C; 8 G; 5 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTT 10
 |||||
 Db 16 CTCTCTCTTT 7

RESULT 70
 ABV82481/c
 ID ABV82481 standard; DNA; 25 BP.

AC ABV82481;

DT 03-JAN-2003 (first entry)

DE Human HTPL scanning oligonucleotide SEQ ID 3727.

XX Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX Homo sapiens.

OS EP1229046-A2.

PN 07-AUG-2002.

PD 28-JAN-2002; 2002EP-0001167.

PF 30-JAN-2001; 2001WO-US000663.

PR 30-JAN-2001; 2001WO-US000664.

PR 30-JAN-2001; 2001WO-US000665.

PR 30-JAN-2001; 2001WO-US000667.

PR 30-JAN-2001; 2001WO-US000668.

PR 23-MAY-2001; 2001US-0864761.

PR 09-OCT-2001; 2001US-0327898.

XX (AEOM-) AEOMICA INC.

XX Zhan J;

XX WPI; 2002-676582/73.

XX Novel isolated human testis expressed Patched like protein (HTPL),
 PT useful for identifying agonist and antagonist and specific binding
 PT partners, and for treating subjects having defects in HTPL -

XX

PS Example 2; Page 552; 718pp; English.

XX The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
 CC shares an overall structure organisation with the Patched protein. The
 CC shared structural features strongly imply that HTPL plays a role similar
 CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC such disorder associated with decreased expression or activity of human
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.

SQ Sequence 25 BP; 9 A; 3 C; 7 G; 6 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;

Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTT 10

|||||
 Db 15 CTCTCTCTTT 6

RESULT 71

ABV82482/c

ID ABV82482 standard; DNA; 25 BP.

XX ABV82482;

XX 03-JAN-2003 (first entry)

DE Human HTPL scanning oligonucleotide SEQ ID 3728.

XX Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.

XX Homo sapiens.

OS EP1229046-A2.

PN 07-AUG-2002.

PD 28-JAN-2002; 2002EP-0001167.

PF 30-JAN-2001; 2001WO-US000663.

PR 30-JAN-2001; 2001WO-US000664.

PR 30-JAN-2001; 2001WO-US000665.

PR 30-JAN-2001; 2001WO-US000667.

PR 30-JAN-2001; 2001WO-US000668.

PR 23-MAY-2001; 2001US-0864761.

PR 09-OCT-2001; 2001US-0327898.

XX (AEOM-) AEOMICA INC.

XX Zhan J;

XX WPI; 2002-676582/73.

XX Novel isolated human testis expressed Patched like protein (HTPL),
 PT useful for identifying agonist and antagonist and specific binding

PT partners, and for treating subjects having defects in HTPL -
 XX
 PS Example 2; Page 552; 718pp; English.
 XX
 CC The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
 CC shares an overall structure organisation with the Patched protein. The
 CC shared structural features strongly imply that HTPL plays a role similar
 CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC therapy and manufacture of a medicament for treatment or prevention of
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.
 XX
 SQ Sequence 25 BP; 9 A; 3 C; 7 G; 6 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 14 CTTCTCTTTT 5
 RESULT 72
 ABV82483/c
 ID ABV82483 standard; DNA; 25 BP.
 AC ABV82483;
 XX
 DT 03-JAN-2003 (first entry)
 XX
 DE Human HTPL scanning oligonucleotide SEQ ID 3729.
 XX
 KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
 XX
 OS Homo sapiens.
 XX
 PN EPI229046-A2.
 XX
 PD 07-AUG-2002.
 XX
 PF 28-JAN-2002; 2002EP-0001167.
 XX
 PR 30-JAN-2001; 2001WO-US00663.
 PR 30-JAN-2001; 2001WO-US00664.
 PR 30-JAN-2001; 2001WO-US00665.
 PR 30-JAN-2001; 2001WO-US00667.
 PR 30-JAN-2001; 2001WO-US00668.
 PR 30-JAN-2001; 2001WO-US00669.
 PR 23-MAY-2001; 2001US-0864761.
 PR 09-OCT-2001; 2001US-0327898.
 XX
 FA (AEOM-) AEOMICA INC.
 XX
 PI Zhan J;
 XX
 DR WPI; 2002-676582/73.
 XX

PT Novel isolated human testis expressed Patched like protein (HTPL),
 PT useful for identifying agonist and antagonist and specific binding
 PT partners, and for treating subjects having defects in HTPL -
 XX
 PS Example 2; Page 552; 718pp; English.
 XX
 CC The present invention relates to human testis expressed Patched like
 CC protein (HTPL, see ABV78759 to ABV78762 and ABB98519 to ABB98520). HTPL
 CC has two isoforms, with a few single base pair differences between the
 CC two. One of the single base pair changes introduces a premature stop
 CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
 CC shares an overall structure organisation with the Patched protein. The
 CC shared structural features strongly imply that HTPL plays a role similar
 CC to that of Patched, and is a potential tumour suppressor. HTPL is
 CC important in regulating male germ cell development, and the HTPL gene was
 CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
 CC useful for diagnosing a disorder caused by mutation in HTPL, and in
 CC therapy and manufacture of a medicament for treatment or prevention of
 CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
 CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
 CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
 CC clinically useful diagnostic markers and potential therapeutic agents for
 CC male infertility and cancer. The present oligonucleotide was used in an
 CC example from the invention.
 XX
 SQ Sequence 25 BP; 9 A; 3 C; 7 G; 6 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 Db 13 CTTCTCTTTT 4
 RESULT 73
 ABV82484/c
 ID ABV82484 standard; DNA; 25 BP.
 AC ABV82484;
 XX
 DT 03-JAN-2003 (first entry)
 XX
 DE Human HTPL scanning oligonucleotide SEQ ID 3730.
 XX
 KW Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
 KW human testis expressed Patched like protein; testis; adrenal; liver;
 KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
 KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
 XX
 OS Homo sapiens.
 XX
 PN EPI229046-A2.
 XX
 PD 07-AUG-2002.
 XX
 PF 28-JAN-2002; 2002EP-0001167.
 XX
 PR 30-JAN-2001; 2001WO-US00663.
 PR 30-JAN-2001; 2001WO-US00664.
 PR 30-JAN-2001; 2001WO-US00665.
 PR 30-JAN-2001; 2001WO-US00667.
 PR 30-JAN-2001; 2001WO-US00668.
 PR 30-JAN-2001; 2001WO-US00669.
 PR 23-MAY-2001; 2001US-0864761.
 PR 09-OCT-2001; 2001US-0327898.
 XX
 FA (AEOM-) AEOMICA INC.
 XX
 PI Zhan J;
 XX
 DR WPI; 2002-676582/73.
 XX

```

DR WPI; 2002-676582/73.
XX
XX Novel isolated human testis expressed Patched like protein (HTPL),
PT useful for identifying agonist and antagonist and specific binding
PT partners, and for treating subjects having defects in HTPL -
XX
XX Example 2; Page 552; 718pp; English.
XX
XX The present invention relates to human testis expressed Patched like
CC protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL
CC has two isoforms, with a few single base pair differences between the
CC two. One of the single base pair changes introduces a premature stop
CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
CC shares an overall structure organisation with the Patched protein. The
CC shared structural features strongly imply that HTPL plays a role similar
CC to that of Patched, and is a potential tumour suppressor. HTPL is
CC important in regulating male germ cell development, and the HTPL gene was
CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
CC useful for diagnosing a disorder caused by mutation in HTPL, and in
CC therapy and manufacture of a medicament for treatment or prevention of
CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
CC clinically useful diagnostic markers and potential therapeutic agents for
CC male infertility and cancer. The present oligonucleotide was used in an
CC example from the invention.
XX
XX Sequence 25 BP; 9 A; 3 C; 7 G; 6 T; 0 other;
SQ
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 12 CTTCTCTTTT 3

RESULT 74
ABV82485/c
ID ABV82485 standard; DNA; 25 BP.
AC ABV82485;
XX
XX 03-JAN-2003 (first entry)
XX
XX Human HTPL scanning oligonucleotide SEQ ID 3731.
XX
XX Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
KW human testis expressed Patched like protein; testis; adrenal; liver;
KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
XX
XX Homo sapiens.
XX
XX EP1229046-A2.
XX
XX 07-AUG-2002.
XX
XX 28-JAN-2002; 2002EP-0001167.
XX
XX 30-JAN-2001; 2001WO-US00663.
XX
XX 30-JAN-2001; 2001WO-US00664.
XX
XX 30-JAN-2001; 2001WO-US00665.
XX
XX 30-JAN-2001; 2001WO-US00667.
XX
XX 30-JAN-2001; 2001WO-US00668.
XX
XX 30-JAN-2001; 2001WO-US00669.
XX
XX 23-MAY-2001; 2001US-0864761.
XX
XX 09-OCT-2001; 2001US-0327898.
XX
XX (AEOM-) AEOMICA INC.
XX
PI Zhan J;
XX
XX WPI; 2002-676582/73.
XX
XX Novel isolated human testis expressed Patched like protein (HTPL),
PT useful for identifying agonist and antagonist and specific binding
PT partners, and for treating subjects having defects in HTPL -
XX
XX Example 2; Page 553; 718pp; English.
XX
XX The present invention relates to human testis expressed Patched like
CC protein (HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL
CC has two isoforms, with a few single base pair differences between the
CC two. One of the single base pair changes introduces a premature stop
CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
CC shares an overall structure organisation with the Patched protein. The
CC shared structural features strongly imply that HTPL plays a role similar
CC to that of Patched, and is a potential tumour suppressor. HTPL is
CC important in regulating male germ cell development, and the HTPL gene was
CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
CC useful for diagnosing a disorder caused by mutation in HTPL, and in
CC therapy and manufacture of a medicament for treatment or prevention of
CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
CC clinically useful diagnostic markers and potential therapeutic agents for
CC male infertility and cancer. The present oligonucleotide was used in an
CC example from the invention.
XX
XX Sequence 25 BP; 9 A; 4 C; 6 G; 6 T; 0 other;
SQ
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 11 CTTCTCTTTT 2

RESULT 75
ABV82486/c
ID ABV82486 standard; DNA; 25 BP.
AC ABV82486;
XX
XX 03-JAN-2003 (first entry)
XX
XX Human HTPL scanning oligonucleotide SEQ ID 3732.
XX
XX Human; gene therapy; tumour suppressor; HTPL; chromosome 10p12.1;
KW human testis expressed Patched like protein; testis; adrenal; liver;
KW male germ cell development; bone marrow; brain; kidney; lung; placenta;
KW prostate; skeletal muscle; colon; male infertility; cancer; ss.
XX
XX Homo sapiens.
XX
XX EP1229046-A2.
XX
XX 07-AUG-2002.
XX
XX 28-JAN-2002; 2002EP-0001167.
XX
XX 30-JAN-2001; 2001WO-US00663.
XX
XX 30-JAN-2001; 2001WO-US00664.
XX
XX 30-JAN-2001; 2001WO-US00665.
XX
XX 30-JAN-2001; 2001WO-US00667.
XX
XX 30-JAN-2001; 2001WO-US00668.
XX
XX 30-JAN-2001; 2001WO-US00669.
XX
XX 23-MAY-2001; 2001US-0864761.
XX
XX 09-OCT-2001; 2001US-0327898.
XX
XX (AEOM-) AEOMICA INC.
XX

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PA (AEOM-) AEOMICA INC.
XX Zhan J;
XX WPI; 2002-676582/73.
XX Novel isolated human testis expressed Patched like protein (HTPL),
PT useful for identifying agonist and antagonist and specific binding
PT partners, and for treating subjects having defects in HTPL -
XX
XX Example 2; Page 553; 718pp; English.
XX The present invention relates to human testis expressed Patched like
CC protein HTPL, see ABV78759 to ABV78762 and AB98519 to AB98520). HTPL
CC has two isoforms, with a few single base pair differences between the
CC two. One of the single base pair changes introduces a premature stop
CC codon in HTPL-S (S for short) compared to HTPL-L (L for long). HTPL
CC shares an overall structure organisation with the Patched protein. The
CC shared structural features strongly imply that HTPL plays a role similar
CC to that of Patched, and is a potential tumour suppressor. HTPL is
CC important in regulating male germ cell development, and the HTPL gene was
CC mapped to human chromosome 10p12.1. HTPL and its coding sequence are
CC useful for diagnosing a disorder caused by mutation in HTPL, and in
CC therapy and manufacture of a medicament for treatment or prevention of
CC such disorder associated with decreased expression or activity of human
CC HTPL. Such disorders include disorders of testis, or adrenal, adult and
CC foetal liver, bone marrow, brain, kidney, lung, placenta, prostate,
CC skeletal muscle or colon function. HTPL proteins and nucleic acids are
CC clinically useful diagnostic markers and potential therapeutic agents for
CC male infertility and cancer. The present oligonucleotide was used in an
CC example from the invention.
XX
XX Sequence 25 BP; 9 A; 4 C; 6 G; 6 T; 0 other;
SQ Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
Db 10 CTTCTCTTTT 1
RESULT 76
ABS75576/C
ID ABS75576 standard; DNA; 25 BP.
AC ABS75576;
XX 27-DEC-2002 (first entry)
XX Human PAPP-Ea associated 25-mer SEQ ID 1102.
DE
DE PAPP-E; human; pregnancy associated plasma protein E; abortive;
KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
KW dysgenetic pregnancy; primer; ss.
XX Homo sapiens.
XX OS
XX US2002102252-A1.
XX 01-AUG-2002.
XX 06-APR-2001; 2001US-0827998.
XX 26-MAY-2000; 2000US-207456P.
XX (GUY/) GU Y.
XX (SHAN/) SHANNON M E.
XX Gu Y, Shannon ME;
XX WPI; 2002-697817/75.
XX New isolated nucleic acid encoding an isoform of human pregnancy
PT associated plasma protein E, for preventing or aborting pregnancy -
XX Example 2; Page 220; 353pp; English.
XX This invention describes a novel isolated nucleic acid that encodes
CC one of three new isoforms of human pregnancy associated plasma protein E,
CC hPAPP-E. The products of the invention have abortive and contraceptive
CC activity and can be used for gene therapy or in a vaccine. The nucleic
CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
CC used in pharmaceutical compositions or vaccines for preventing or
CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
CC antibodies can be used to assess the expression levels of PAPP-E isoform
CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
CC antenatally. This sequence represents an oligomer used in scanning the
CC human PAPP-E genes described in the disclosure of the invention.
XX
XX Sequence 25 BP; 11 A; 3 C; 8 G; 3 T; 0 other;
SQ Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16
RESULT 77
ABS75577/C
ID ABS75577 standard; DNA; 25 BP.
AC ABS75577;
XX 27-DEC-2002 (first entry)
XX Human PAPP-Ea associated 25-mer SEQ ID 1103.
DE
DE PAPP-E; human; pregnancy associated plasma protein E; abortive;
KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
KW dysgenetic pregnancy; primer; ss.
XX Homo sapiens.
XX OS
XX US2002102252-A1.
XX 01-AUG-2002.
XX 06-APR-2001; 2001US-0827998.
XX 26-MAY-2000; 2000US-207456P.
XX (GUY/) GU Y.
XX (SHAN/) SHANNON M E.
XX Gu Y, Shannon ME;
XX WPI; 2002-697817/75.
XX New isolated nucleic acid encoding an isoform of human pregnancy
PT associated plasma protein E, for preventing or aborting pregnancy -
XX Example 2; Page 220; 353pp; English.
XX This invention describes a novel isolated nucleic acid that encodes
CC one of three new isoforms of human pregnancy associated plasma protein E,
CC hPAPP-E. The products of the invention have abortive and contraceptive
CC activity and can be used for gene therapy or in a vaccine. The nucleic
CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
CC used in pharmaceutical compositions or vaccines for preventing or

CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 CC antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.

XX
 SQ Sequence 25 BP; 11 A; 3 C; 8 G; 3 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||
 Db 24 CTTCTCTTTT 15

RESULT 78
 ABS75578/c
 ID ABS75578 standard; DNA; 25 BP.
 AC ABS75578;
 XX
 XX
 DT 27-DEC-2002 (first entry)
 XX
 DE Human PAPP-Ea associated 25-mer SEQ ID 1104.
 XX
 KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
 KW dysgenetic pregnancy; primer; ss.
 XX
 OS Homo sapiens.
 XX
 PN US2002102252-A1.
 XX
 PD 01-AUG-2002.
 XX
 PF 06-APR-2001; 2001US-0827998.
 XX
 PR 26-MAY-2000; 2000US-207456P.
 XX
 PA (GUY/) GU Y.
 PA (SHAN/) SHANNON M E.
 XX
 PI Gu Y, Shannon ME;
 XX
 DR WPI; 2002-697817/75.
 XX
 PT New isolated nucleic acid encoding an isoform of human pregnancy
 PT associated plasma protein E, for preventing or aborting pregnancy -
 XX
 PS Example 2; Page 220; 353pp; English.

XX This invention describes a novel isolated nucleic acid that encodes
 CC one of three new isoforms of human pregnancy associated plasma protein E,
 CC hPAPP-E. The products of the invention have abortive and contraceptive
 CC activity and can be used for gene therapy or in a vaccine. The nucleic
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
 CC used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 CC antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.

XX
 SQ Sequence 25 BP; 11 A; 3 C; 7 G; 4 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 CTTCTCTTTT 10
 |||||
 Db 23 CTTCTCTTTT 14

RESULT 79
 ABS75579/c
 ID ABS75579 standard; DNA; 25 BP.
 XX
 AC ABS75579;
 XX
 DT 27-DEC-2002 (first entry)
 XX
 DE Human PAPP-Ea associated 25-mer SEQ ID 1105.
 XX
 KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
 KW dysgenetic pregnancy; primer; ss.
 XX
 OS Homo sapiens.
 XX
 PN US2002102252-A1.
 XX
 PD 01-AUG-2002.
 XX
 PF 06-APR-2001; 2001US-0827998.
 XX
 PR 26-MAY-2000; 2000US-207456P.
 XX
 PA (GUY/) GU Y.
 PA (SHAN/) SHANNON M E.
 XX
 PI Gu Y, Shannon ME;
 XX
 DR WPI; 2002-697817/75.
 XX
 PT New isolated nucleic acid encoding an isoform of human pregnancy
 PT associated plasma protein E, for preventing or aborting pregnancy -
 XX
 PS Example 2; Page 220; 353pp; English.

XX This invention describes a novel isolated nucleic acid that encodes
 CC one of three new isoforms of human pregnancy associated plasma protein E,
 CC hPAPP-E. The products of the invention have abortive and contraceptive
 CC activity and can be used for gene therapy or in a vaccine. The nucleic
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
 CC used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 CC antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.

XX
 SQ Sequence 25 BP; 12 A; 2 C; 7 G; 4 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||
 Db 22 CTTCTCTTTT 13

RESULT 80
 ABS75580/c
 ID ABS75580 standard; DNA; 25 BP.
 XX
 AC ABS75580;


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XX 27-DEC-2002 (first entry)
XX DT
XX DE
XX DE Human PAPP-Ea associated 25-mer SEQ ID 1106.
XX
XX PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX KW dysgenetic pregnancy; primer; ss.
XX
XX OS Homo sapiens.
XX
XX US2002102252-A1.
XX PN
XX
XX 01-AUG-2002.
XX PD
XX PF
XX PF 06-APR-2001; 2001US-0827998.
XX
XX 26-MAY-2000; 2000US-207456P.
XX PR
XX
XX (GUY/) GU Y.
XX PA (SHAN/) SHANNON M E.
XX
XX PI Gu Y, Shannon ME;
XX
XX WPI; 2002-697817/75.
XX
XX New isolated nucleic acid encoding an isoform of human pregnancy
XX associated plasma protein E, for preventing or aborting pregnancy
XX
XX Example 2; Page 220; 353pp; English.
XX
XX This invention describes a novel isolated nucleic acid that encodes
XX one of three new isoforms of human pregnancy associated plasma protein E,
XX hPAPP-E. The products of the invention have abortive and contraceptive
XX activity and can be used for gene therapy or in a vaccine. The nucleic
XX acid, polypeptide encoded by it, or antibody to the polypeptide can be
XX used in pharmaceutical compositions or vaccines for preventing or
XX aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
XX dysgenetic pregnancies. The nucleic acids are used as probes to assess
XX the level of PAPP-E isoform mRNA in chorionic villus samples, and the
XX antibodies can be used to assess the expression levels of PAPP-E isoform
XX proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
XX antenatally. This sequence represents an oligomer used in scanning the
XX human PAPP-E genes described in the disclosure of the invention.
XX
XX Sequence 25 BP; 13 A; 2 C; 7 G; 3 T; 0 other;
XX SQ
XX
XX Query Match 100.0%; Score 10; DB 24; Length 25;
XX Best Local Similarity 100.0%; Pred. No. 1.8e+04;
XX Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 CTTCTCTTTT 10
XX Db |||||
XX 21 CTTCTCTTTT 12
XX
XX RESULT 81
XX ABS75581/C
XX ID ABS75581 standard; DNA; 25 BP.
XX
XX AC ABS75581;
XX
XX AC ABS75581;
XX
XX 27-DEC-2002 (first entry)
XX DT
XX DE Human PAPP-Ea associated 25-mer SEQ ID 1107.
XX
XX PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX KW dysgenetic pregnancy; primer; ss.
XX
XX OS Homo sapiens.
XX
XX US2002102252-A1.
XX PN

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XX 01-AUG-2002.
XX PD
XX PF
XX PF 06-APR-2001; 2001US-0827998.
XX
XX 26-MAY-2000; 2000US-207456P.
XX PR
XX
XX (GUY/) GU Y.
XX PA (SHAN/) SHANNON M E.
XX
XX PI Gu Y, Shannon ME;
XX
XX WPI; 2002-697817/75.
XX
XX New isolated nucleic acid encoding an isoform of human pregnancy
XX associated plasma protein E, for preventing or aborting pregnancy
XX
XX Example 2; Page 220; 353pp; English.
XX
XX This invention describes a novel isolated nucleic acid that encodes
XX one of three new isoforms of human pregnancy associated plasma protein E,
XX hPAPP-E. The products of the invention have abortive and contraceptive
XX activity and can be used for gene therapy or in a vaccine. The nucleic
XX acid, polypeptide encoded by it, or antibody to the polypeptide can be
XX used in pharmaceutical compositions or vaccines for preventing or
XX aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
XX dysgenetic pregnancies. The nucleic acids are used as probes to assess
XX the level of PAPP-E isoform mRNA in chorionic villus samples, and the
XX antibodies can be used to assess the expression levels of PAPP-E isoform
XX proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
XX antenatally. This sequence represents an oligomer used in scanning the
XX human PAPP-E genes described in the disclosure of the invention.
XX
XX Sequence 25 BP; 13 A; 2 C; 7 G; 3 T; 0 other;
XX SQ
XX
XX Query Match 100.0%; Score 10; DB 24; Length 25;
XX Best Local Similarity 100.0%; Pred. No. 1.8e+04;
XX Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 CTTCTCTTTT 10
XX Db |||||
XX 20 CTTCTCTTTT 11
XX
XX RESULT 82
XX ABS75582/C
XX ID ABS75582 standard; DNA; 25 BP.
XX
XX AC ABS75582;
XX
XX AC ABS75582;
XX
XX 27-DEC-2002 (first entry)
XX DT
XX DE Human PAPP-Ea associated 25-mer SEQ ID 1108.
XX
XX PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX KW dysgenetic pregnancy; primer; ss.
XX
XX OS Homo sapiens.
XX
XX OS US2002102252-A1.
XX PN
XX
XX 01-AUG-2002.
XX PD
XX PF
XX PF 06-APR-2001; 2001US-0827998.
XX
XX 26-MAY-2000; 2000US-207456P.
XX PR
XX
XX (GUY/) GU Y.
XX PA (SHAN/) SHANNON M E.
XX
XX PI Gu Y, Shannon ME;
XX
XX US2002102252-A1.
XX PN

```

DR WPI; 2002-697817/75.
 XX New isolated nucleic acid encoding an isoform of human pregnancy
 PT associated plasma protein E, for preventing or aborting pregnancy
 XX
 PS Example 2; Page 221; 353pp; English.
 XX
 CC This invention describes a novel isolated nucleic acid that encodes
 CC one of three new isoforms of human pregnancy associated plasma protein E,
 CC hPAPP-E. The products of the invention have abortive and contraceptive
 CC activity and can be used for gene therapy or in a vaccine. The nucleic
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
 CC used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 CC antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.
 XX
 SQ Sequence 25 BP; 12 A; 2 C; 7 G; 4 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 CTTCTCTTTT 10
 Db 19 CTTCTCTTTT 10
 RESULT 83
 ABS75583/C
 ID ABS75583 standard; DNA; 25 BP.
 XX
 AC ABS75583;
 XX
 DT 27-DEC-2002 (first entry)
 XX
 DE Human PAPP-Ea associated 25-mer SEQ ID 1109.
 XX
 KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
 KW dysgenetic pregnancy; primer; ss.
 XX
 OS Homo sapiens.
 XX
 PN US2002102252-A1.
 XX
 PD 01-AUG-2002.
 XX
 PF 06-APR-2001; 2001US-0827998.
 XX
 PR 26-MAY-2000; 2000US-207456P.
 XX
 PA (GUY/) GU Y.
 XX (SHAN/) SHANNON M E.
 PI Gu Y, Shannon ME;
 XX WPI; 2002-697817/75.
 XX
 PT New isolated nucleic acid encoding an isoform of human pregnancy
 PT associated plasma protein E, for preventing or aborting pregnancy
 XX
 PS Example 2; Page 221; 353pp; English.
 XX
 CC This invention describes a novel isolated nucleic acid that encodes
 CC one of three new isoforms of human pregnancy associated plasma protein E,
 CC hPAPP-E. The products of the invention have abortive and contraceptive
 CC activity and can be used for gene therapy or in a vaccine. The nucleic
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be

CC used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 CC antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.
 XX
 SQ Sequence 25 BP; 12 A; 3 C; 6 G; 4 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 CTTCTCTTTT 10
 Db 18 CTTCTCTTTT 9
 RESULT 84
 ABS75584/C
 ID ABS75584 standard; DNA; 25 BP.
 XX
 AC ABS75584;
 XX
 DT 27-DEC-2002 (first entry)
 XX
 DE Human PAPP-Ea associated 25-mer SEQ ID 1110.
 XX
 KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
 KW dysgenetic pregnancy; primer; ss.
 XX
 OS Homo sapiens.
 XX
 PN US2002102252-A1.
 XX
 PD 01-AUG-2002.
 XX
 PF 06-APR-2001; 2001US-0827998.
 XX
 PR 26-MAY-2000; 2000US-207456P.
 XX
 PA (GUY/) GU Y.
 XX (SHAN/) SHANNON M E.
 PI Gu Y, Shannon ME;
 XX WPI; 2002-697817/75.
 XX
 PT New isolated nucleic acid encoding an isoform of human pregnancy
 PT associated plasma protein E, for preventing or aborting pregnancy
 XX
 PS Example 2; Page 221; 353pp; English.
 XX
 CC This invention describes a novel isolated nucleic acid that encodes
 CC one of three new isoforms of human pregnancy associated plasma protein E,
 CC hPAPP-E. The products of the invention have abortive and contraceptive
 CC activity and can be used for gene therapy or in a vaccine. The nucleic
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
 CC used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 CC antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.
 XX
 SQ Sequence 25 BP; 12 A; 4 C; 5 G; 4 T; 0 other;
 Query Match 100.0%; Score 10; DB 24; Length 25;

```
Best Local Similarity 100.0%; Pred. No. 1.8e+04; Mismatches 0; Indels 0; Gaps 0;
Matches 10; Conservative 0;

QY 1 CTTCTCTTTT 10
Db 17 CTTCTCTTTT 8

RESULT 85
ABS75585/c
ID ABS75585 standard; DNA; 25 BP.
XX
AC ABS75585;
XX
DT 27-DEC-2002 (first entry)
XX
DE Human PAPP-Ea associated 25-mer SEQ ID 1111.
XX
KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
KW dysgenetic pregnancy; primer; ss.
XX
OS Homo sapiens.
XX
PN US2002102252-A1.
XX
PD 01-AUG-2002.
XX
PF 06-APR-2001; 2001US-0827998.
XX
PR 26-MAY-2000; 2000US-207456P.
XX
PA (GUY/) GU Y.
PA (SHAN/) SHANNON M E.
XX
PI Gu Y, Shannon ME;
XX
PI WPI; 2002-697817/75.
XX
PT New isolated nucleic acid encoding an isoform of human pregnancy
PT associated plasma protein E, for preventing or aborting pregnancy
PS Example 2; Page 221; 353pp; English.
XX
XX This invention describes a novel isolated nucleic acid that encodes
XX one of three new isoforms of human pregnancy associated plasma protein E,
XX hPAPP-E. The products of the invention have abortive and contraceptive
XX activity and can be used for gene therapy or in a vaccine. The nucleic
XX acid, polypeptide encoded by it, or antibody to the polypeptide can be
XX used in pharmaceutical compositions or vaccines for preventing or
XX aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
XX dysgenetic pregnancies. The nucleic acids are used as probes to assess
XX the level of PAPP-E isoform mRNA in chorionic villus samples, and the
XX antibodies can be used to assess the expression levels of PAPP-E isoform
XX proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
XX antenatally. This sequence represents an oligomer used in scanning the
XX human PAPP-E genes described in the disclosure of the invention.
XX
SQ Sequence 25 BP; 12 A; 4 C; 4 G; 5 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 16 CTTCTCTTTT 7

RESULT 86
ABS75586/c
ID ABS75586 standard; DNA; 25 BP.
XX
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```
AC ABS75586;
XX
DT 27-DEC-2002 (first entry)
XX
DE Human PAPP-Ea associated 25-mer SEQ ID 1112.
XX
KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
KW dysgenetic pregnancy; primer; ss.
XX
OS Homo sapiens.
XX
PN US2002102252-A1.
XX
PD 01-AUG-2002.
XX
PF 06-APR-2001; 2001US-0827998.
XX
PR 26-MAY-2000; 2000US-207456P.
XX
PA (GUY/) GU Y.
PA (SHAN/) SHANNON M E.
XX
PI Gu Y, Shannon ME;
XX
PI WPI; 2002-697817/75.
XX
PT New isolated nucleic acid encoding an isoform of human pregnancy
PT associated plasma protein E, for preventing or aborting pregnancy
PS Example 2; Page 221; 353pp; English.
XX
XX This invention describes a novel isolated nucleic acid that encodes
XX one of three new isoforms of human pregnancy associated plasma protein E,
XX hPAPP-E. The products of the invention have abortive and contraceptive
XX activity and can be used for gene therapy or in a vaccine. The nucleic
XX acid, polypeptide encoded by it, or antibody to the polypeptide can be
XX used in pharmaceutical compositions or vaccines for preventing or
XX aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
XX dysgenetic pregnancies. The nucleic acids are used as probes to assess
XX the level of PAPP-E isoform mRNA in chorionic villus samples, and the
XX antibodies can be used to assess the expression levels of PAPP-E isoform
XX proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
XX antenatally. This sequence represents an oligomer used in scanning the
XX human PAPP-E genes described in the disclosure of the invention.
XX
SQ Sequence 25 BP; 12 A; 4 C; 4 G; 5 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 15 CTTCTCTTTT 6

RESULT 87
ABS75587/c
ID ABS75587 standard; DNA; 25 BP.
XX
AC ABS75587;
XX
DT 27-DEC-2002 (first entry)
XX
DE Human PAPP-Ea associated 25-mer SEQ ID 1113.
XX
KW PAPP-E; human; pregnancy associated plasma protein E; abortive;
KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
KW dysgenetic pregnancy; primer; ss.
XX
OS Homo sapiens.
XX
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PN US2002102252-A1.
XX
XX 01-AUG-2002.
XX
XX 06-APR-2001; 2001US-0827998.
XX
XX 26-MAY-2000; 2000US-207456P.
XX
XX (GUY/ ) GU Y.
PA (SHAN/ ) SHANNON M E.
XX
XX Gu Y, Shannon ME;
XX
XX WPI; 2002-697817/75.
XX
XX New isolated nucleic acid encoding an isoform of human pregnancy
PT associated plasma protein E, for preventing or aborting pregnancy
XX
XX Example 2; Page 221; 353pp; English.
XX
XX This invention describes a novel isolated nucleic acid that encodes
CC one of three new isoforms of human pregnancy associated plasma protein E,
CC hPAPP-E. The products of the invention have abortive and contraceptive
CC activity and can be used for gene therapy or in a vaccine. The nucleic
CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
CC used in pharmaceutical compositions or vaccines for preventing or
CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
CC antibodies can be used to assess the expression levels of PAPP-E isoform
CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
CC antenatally. This sequence represents an oligomer used in scanning the
CC human PAPP-E genes described in the disclosure of the invention.
XX
XX Sequence 25 BP; 12 A; 3 C; 5 G; 5 T; 0 other;
SQ
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 14 CTTCTCTTTT 5

RESULT 88
ABS75588/c
ID ABS75588 standard; DNA; 25 BP.
XX
XX ABS75588;
XX
XX 27-DEC-2002 (first entry)
XX
XX Human PAPP-Ea associated 25-mer SEQ ID 1114.
XX
XX PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX dysgenetic pregnancy; primer; ss.
XX
XX Homo sapiens.
XX
XX US2002102252-A1.
XX
XX 01-AUG-2002.
XX
XX 06-APR-2001; 2001US-0827998.
XX
XX 26-MAY-2000; 2000US-207456P.
XX
XX (GUY/ ) GU Y.
XX (SHAN/ ) SHANNON M E.
XX
XX Gu Y, Shannon ME;
XX
XX WPI; 2002-697817/75.
XX
XX New isolated nucleic acid encoding an isoform of human pregnancy
PT associated plasma protein E, for preventing or aborting pregnancy
XX
XX Example 2; Page 221; 353pp; English.
XX
XX This invention describes a novel isolated nucleic acid that encodes
CC one of three new isoforms of human pregnancy associated plasma protein E,
CC hPAPP-E. The products of the invention have abortive and contraceptive
CC activity and can be used for gene therapy or in a vaccine. The nucleic
CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
CC used in pharmaceutical compositions or vaccines for preventing or
CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
CC antibodies can be used to assess the expression levels of PAPP-E isoform
CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
CC antenatally. This sequence represents an oligomer used in scanning the
CC human PAPP-E genes described in the disclosure of the invention.
XX
XX Sequence 25 BP; 12 A; 3 C; 5 G; 5 T; 0 other;
SQ
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

RESULT 89
ABS75589/c
ID ABS75589 standard; DNA; 25 BP.
XX
XX ABS75589;
XX
XX 27-DEC-2002 (first entry)
XX
XX Human PAPP-Ea associated 25-mer SEQ ID 1115.
XX
XX PAPP-E; human; pregnancy associated plasma protein E; abortive;
XX contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
XX dysgenetic pregnancy; primer; ss.
XX
XX Homo sapiens.
XX
XX US2002102252-A1.
XX
XX 01-AUG-2002.
XX
XX 06-APR-2001; 2001US-0827998.
XX
XX 26-MAY-2000; 2000US-207456P.
XX
XX (GUY/ ) GU Y.
XX (SHAN/ ) SHANNON M E.
XX
XX Gu Y, Shannon ME;
XX
XX WPI; 2002-697817/75.
XX
XX New isolated nucleic acid encoding an isoform of human pregnancy
PT associated plasma protein E, for preventing or aborting pregnancy
XX
XX Example 2; Page 221; 353pp; English.
XX
XX This invention describes a novel isolated nucleic acid that encodes
CC one of three new isoforms of human pregnancy associated plasma protein E,
CC hPAPP-E. The products of the invention have abortive and contraceptive
CC activity and can be used for gene therapy or in a vaccine. The nucleic
CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
CC used in pharmaceutical compositions or vaccines for preventing or
CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
CC antibodies can be used to assess the expression levels of PAPP-E isoform
CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
CC antenatally. This sequence represents an oligomer used in scanning the
CC human PAPP-E genes described in the disclosure of the invention.
XX
XX Sequence 25 BP; 12 A; 3 C; 6 G; 4 T; 0 other;
SQ
Query Match 100.0%; Score 10; DB 24; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.8e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

```

CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
 CC used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 CC antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.
 XX
 SQ Sequence 25 BP; 11 A; 3 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTTT 10
 |||||
 Db 12 CTCTCTCTTTT 3

RESULT 90
 ABS75590/c
 ID ABS75590 standard; DNA; 25 BP.

XX AC ABS75590;
 XX DT 27-DEC-2002 (first entry)
 XX DE Human PAPP-Ea associated 25-mer SEQ ID 1116.

XX PAPP-E; human; pregnancy associated plasma protein E; abortive;
 XX contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
 KW dysgenetic pregnancy; primer; ss.
 XX OS Homo sapiens.

XX PN US2002102252-A1.
 XX PD 01-AUG-2002.

XX PF 06-APR-2001; 2001US-0827998.
 XX PR 26-MAY-2000; 2000US-207456P.

XX PA (GUY/) GU Y.
 XX PA (SHAN/) SHANNON M E.

XX PI Gu Y, Shannon ME;
 XX DR WPI; 2002-697817/75.

XX PT New isolated nucleic acid encoding an isoform of human pregnancy
 PT associated plasma protein E, for preventing or aborting pregnancy

XX FS Example 2; Page 222; 353pp; English.

XX This invention describes a novel isolated nucleic acid that encodes
 CC one of three new isoforms of human pregnancy associated plasma protein E,
 CC hPAPP-E. The products of the invention have abortive and contraceptive
 CC activity and can be used for gene therapy or in a vaccine. The nucleic
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
 CC used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 CC antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.

XX SQ Sequence 25 BP; 12 A; 2 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTTT 10
 |||||
 Db 11 CTCTCTCTTTT 2

RESULT 91
 ABS75591/c
 ID ABS75591 standard; DNA; 25 BP.

XX AC ABS75591;
 XX DT 27-DEC-2002 (first entry)

XX DE Human PAPP-Ea associated 25-mer SEQ ID 1117.
 XX PAPP-E; human; pregnancy associated plasma protein E; abortive;
 KW contraceptive; gene therapy; vaccine; pregnancy; antenatal; diagnosis;
 KW dysgenetic pregnancy; primer; ss.

XX OS Homo sapiens.
 XX PN US2002102252-A1.

XX PD 01-AUG-2002.

XX PF 06-APR-2001; 2001US-0827998.

XX PR 26-MAY-2000; 2000US-207456P.

XX PA (GUY/) GU Y.
 XX PA (SHAN/) SHANNON M E.

XX PI Gu Y, Shannon ME;
 XX DR WPI; 2002-697817/75.

XX PT New isolated nucleic acid encoding an isoform of human pregnancy
 PT associated plasma protein E, for preventing or aborting pregnancy

XX FS Example 2; Page 222; 353pp; English.

XX This invention describes a novel isolated nucleic acid that encodes
 CC one of three new isoforms of human pregnancy associated plasma protein E,
 CC hPAPP-E. The products of the invention have abortive and contraceptive
 CC activity and can be used for gene therapy or in a vaccine. The nucleic
 CC acid, polypeptide encoded by it, or antibody to the polypeptide can be
 CC used in pharmaceutical compositions or vaccines for preventing or
 CC aborting pregnancy. PAPP-E is used in the antenatal diagnosis of
 CC dysgenetic pregnancies. The nucleic acids are used as probes to assess
 CC the level of PAPP-E isoform mRNA in chorionic villus samples, and the
 CC antibodies can be used to assess the expression levels of PAPP-E isoform
 CC proteins in chorionic villus samples, to diagnose dysgenetic pregnancies
 CC antenatally. This sequence represents an oligomer used in scanning the
 CC human PAPP-E genes described in the disclosure of the invention.

XX SQ Sequence 25 BP; 12 A; 2 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 24; Length 25;
 Best Local Similarity 100.0%; Pred. No. 1.8e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTTT 10
 |||||
 Db 10 CTCTCTCTTTT 1

RESULT 92
 AAQ45398/c
 ID AAQ45398 standard; DNA; 27 BP.

XX AC AAQ45398;
 XX XX
 DT 25-MAR-2003 (updated)
 DT 11-NOV-1994 (first entry)
 XX XX
 DE Oligonucleotide forming triplex with viral polypurine tract.
 XX XX
 KW HIV; human immunodeficiency virus; retrovirus; hepatitis virus;
 KW reverse transcription; virus replication; inhibition; treatment;
 KW therapy; polypurine; triplex; antisense; ss.
 XX OS Synthetic.
 XX XX
 FN WO9407367-A1.
 XX XX
 PD 14-APR-1994.
 XX XX
 PF 29-SEP-1993; 93WO-US09300.
 XX XX
 PR 29-SEP-1992; 92US-0954184.
 XX XX
 PA (APOL-) APOLLON INC.
 PA (PLAC) MAX PLANCK GES FOERDERUNG WISSENSCHAFTEN.
 XX XX
 PI Moelling K;
 XX XX
 DR WPI; 1994-135099/16.
 XX XX
 PS Antiviral oligomers that bind poly-purine tracts of
 PT single-stranded RNA or RNA - DNA hybrids - used to target the
 PT early stages of viral replication before double stranded DNA is
 PT formed
 XX XX
 XX Example 4; Page 29; 54pp; English.
 XX XX
 CC Administration of antisense or triplex forming oligonucleotides
 CC which bind polypurine tracts (PPT) may be used in the therapy or
 CC treatment of individuals infected with retroviruses or hepatitis
 CC viruses since in these two families of viruses, two primers are
 CC involved in the reverse transcription of viral RNA into double
 CC stranded DNA, one of which is a PPT. The antisense or triplex
 CC forming oligonucleotides can inhibit the early stages of viral
 CC replication by binding to the PPT primer or by binding to PPT
 CC tracts in the RNA-DNA hybrid molecule formed after reverse
 CC transcription of the viral RNA. This oligonucleotide was incubated
 CC with an in vitro transcribed 5' end labelled pKJ2 RNA of 134
 CC nucleotides in length to which a 40-mer deoxyribonucleotide
 CC complementary to the PPT had been hybridised. The presence of this
 CC sequence led to protection of the pKJ2 RNA-DNA hybrid from RNase H
 CC digestion which would suggest the formation of a triplex. Triplex
 CC formation was confirmed using a primer extension technique. A
 CC primer binding downstream of the PPT was synthesised and extended in
 CC vitro by reverse transcriptase in the presence of
 CC oligodeoxynucleotides including one which was radioactively labelled.
 CC The newly synthesised DNA was terminated at the site of the PPT when
 CC the triplex was formed and blocked extension. Triplex formation
 CC would be expected to interfere with viral replication in vivo.
 CC See AAQ45381-Q45417. This is a variant of the sequence described in
 CC AAQ45387.
 CC (Updated on 25-MAR-2003 to correct PN field.)
 XX XX
 SQ Sequence 27 BP; 7 A; 8 C; 8 G; 4 T; 0 other;

Query Match 100.0%; Score 10; DB 15; Length 27;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 |||||
 DB 22 CTTCTCTTTT 13

RESULT 93
 AAX74299/c
 ID AAX74299 standard; RNA; 27 BP.
 XX XX
 AC AAX74299;
 XX XX
 DT 28-JUL-1999 (first entry)
 XX XX
 DE Mouse flt-1 VEGF receptor hammerhead ribozyme #771.
 XX XX
 KW Vascular endothelial growth factor receptor; VEGF receptor; flt-1;
 KW flk-1; KDR; hammerhead ribozyme; hairpin ribozyme; cleavage;
 KW tumour angiogenesis; psoriasis; rheumatoid arthritis; ocular disease;
 KW fms-like tyrosine kinase 1; kinase insert domain containing receptor;
 KW foetal liver kinase 1; ss.
 XX XX
 OS Synthetic.
 OS Mus sp.
 XX XX
 FN WO9715662-A2.
 XX XX
 PD 01-MAY-1997.
 XX XX
 PF 25-OCT-1996; 96WO-US17480.
 XX XX
 PR 11-JAN-1996; 96US-0584040.
 PR 26-OCT-1995; 95US-0005974.
 XX XX
 PA (CHIR) CHIRON CORP.
 PA (RIBO-) RIBOZYME PHARM INC.
 XX XX
 PI Escobedo J, McSwiggen J, Pavco P, Stinchcomb D;
 XX XX
 DR WPI; 1997-259017/23.
 XX XX
 PT Nucleic acid molecule modulating VEGF receptor(s) gene expression or
 PT mRNA stability - useful for treating e.g. tumour angiogenesis,
 PT psoriasis, rheumatoid arthritis, etc., in a human patient
 XX XX
 PS Claim 9; Page 178; 218pp; English.
 XX XX
 CC The present invention describes nucleic acid molecules which modulate
 CC the synthesis, expression and/or stability of a mRNA encoding 1 or more
 CC receptors of vascular endothelial growth factor (VEGF). A patient
 CC (preferably human) having a condition associated with the level of the
 CC fms-like tyrosine kinase 1 (flt-1), kinase insert domain containing
 CC receptor (KDR) and/or foetal liver kinase 1 (flk-1) (e.g. tumour
 CC angiogenesis, ocular diseases, psoriasis and rheumatoid arthritis) can
 CC be treated by administering the nucleic acid molecule or the expression
 CC vector to the patient. AAX67275 to AAX75752 represent specific examples
 CC of nucleic acid molecules from the present invention.
 XX XX
 SQ Sequence 27 BP; 12 A; 1 C; 9 G; 4 U; 1 other;
 Query Match 100.0%; Score 10; DB 18; Length 27;
 Best Local Similarity 100.0%; Pred. No. 1.7e+04;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 |||||
 DB 27 CTTCTCTTTT 18

RESULT 94
 AAH62997
 ID AAH62997 standard; DNA; 27 BP.
 XX XX
 AC AAH62997;
 XX XX
 DT 11-SEP-2001 (first entry)
 XX XX
 DE Shrimp white spot Bacilliform virus (WSBV) oligonucleotide 158.
 XX XX

KW Shrimp white spot Bacilliform virus; WSBV; diagnosis; viral infection;
KW antiviral agent; gene expression; antisense construct; probe; primer;
KW transgenic viral resistant shrimp; ss.
XX
XX
OS White spot syndrome virus.
XX
XX WO200138351-A2.
XX
XX 31-MAY-2001.
XX
XX 08-NOV-2000; 2000WO-US28888.
XX
XX 24-NOV-1999; 99CN-0124717.
XX
XX (PENY-) PE CORP NY.
XX
XX PA THIRD INST OCEANOGRAPHY STATE OCEANI C A.
XX
XX (SINO-) SINOGENOMAX CO LTD.
XX
XX Xu X, Yang F, He J, Pham L, He M, Ye Y, Shen Y, Kodira C;
XX
XX WPI; 2001-355877/37.
XX
XX Primary nucleotide sequence of the shrimp white spot Bacilliform virus
XX (WSBV), useful for producing viral polypeptides that can be used to
XX screen for agents that are useful for treating WSBV infection -
XX
XX Disclosure; Figure 3; 626pp; English.
XX
XX The invention provides the primary nucleotide sequence of the WSBV genome
XX (AAH62689), predicted transcript sequences (AAH62689-AAH62839) and
XX encoded proteins (AAH62840-AAH62851) and oligonucleotide sequences
XX (AAH62840-63160) suitable for use as primers or probes. The nucleic acid
XX molecules and proteins of the invention are useful for diagnosis and
XX monitoring viral infection, in screens for antiviral agents and for
XX monitoring viral gene expression or activity during a treatment regimen.
XX The nucleic acid molecules are also useful as antisense constructs to
XX control viral gene expression in infected cells and tissues and to create
XX transgenic viral resistant shrimp.
XX
SQ Sequence 27 BP; 5 A; 6 C; 2 G; 14 T; 0 other;

Query Match 100.0%; Score 10; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 2 CTTCTCTTTT 11

RESULT 95
AAH38166/C
ID AAH38166 standard; DNA; 27 BP.
XX
XX AAH38166;
XX
XX 14-AUG-2001 (first entry)
XX
XX SNP specific lower PCR primer SEQ ID 962.
XX
XX Single nucleotide polymorphism; SNP; single nucleotide primer extension;
XX SNPE; genotyping; agammaglobulinaemia; diabetes insipidus; cancer;
XX Lesch-Nyhan syndrome; muscular dystrophy; familial hypercholesterolaemia;
XX polycystic kidney disease; osteogenesis imperfecta; autoimmune disease;
XX acute intermittent porphyria; rheumatoid arthritis; multiple sclerosis;
XX inflammation; forensic investigation; paternity analysis; PCR primer; ss.
XX
XX Homo sapiens.
XX
XX WO200129262-A2.
XX
XX 26-APR-2001.
XX

PF 13-OCT-2000; 2000WO-US28436.
XX
XX 15-OCT-1999; 99US-0160096.
XX
XX (ORCH-) ORCHID BIOSCIENCES INC.
XX
XX Picoult-Newburg L, Pohl M;
XX
XX WPI; 2001-290930/30.
XX
XX New genotyping oligonucleotide, useful for detecting the presence,
XX PT absence or identity of single polynucleotide polymorphism in a nucleic
XX PT acid sample -
XX
XX Claim 1; Page 54; 83pp; English.
XX
XX Sequences AAH37205 - AAH40944 represent PCR primers, single nucleotide
XX primer extension (SNPE) primers, and the sequences of regions flanking
XX sites of single nucleotide polymorphisms SNPs. The present invention
XX includes kits for determining the presence or absence of a SNP, using the
XX oligonucleotides of the invention. The PCR primers are used to amplify a
XX SNP flanking sequence, the SNPE primer is used as a genotyping primer.
XX The oligonucleotides are useful for genotyping a nucleic acid sample by
XX performing a single-nucleotide primer extension reaction. The
XX oligonucleotides are useful for determining the presence, absence or
XX identity of a SNP and for genotyping nucleic acid samples, for e.g. to
XX assess by association analysis the genotype of an individual or group of
XX individuals, having a pathological phenotypic trait suspected of being
XX caused by one or more SNPs. Phenotypic traits include diseases e.g.
XX agammaglobulinaemia, diabetes insipidus, Lesch-Nyhan syndrome, muscular
XX dystrophy, familial hypercholesterolaemia, polycystic kidney disease,
XX osteogenesis imperfecta and acute intermittent porphyria. Phenotypic
XX traits also include symptoms of or susceptibility to multifactorial
XX disease of which a component is or may be genetic such as autoimmune
XX diseases, including, rheumatoid arthritis, multiple sclerosis,
XX inflammation, cancer, nervous system diseases and infection by pathogenic
XX microorganism. The method is also useful in forensic investigations and
XX paternity analysis. The present sequence represents a PCR primer specific
XX for a human SNP containing DNA sequence.
XX
SQ Sequence 27 BP; 13 A; 5 C; 4 G; 5 T; 0 other;

Query Match 100.0%; Score 10; DB 22; Length 27;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 21 CTTCTCTTTT 12

RESULT 96
AAC81508
ID AAC81508 standard; DNA; 28 BP.
XX
XX AAC81508;
XX
XX 28-FEB-2001 (first entry)
XX
XX Human MSF exon 6 RT-PCR primer, SEQ ID NO:16.
XX
XX Human MSF; megakaryocyte stimulating factor; tribonectin;
XX alternative splicing; joint boundary lubricant; O-linked oligosaccharide;
XX osteoarthritis; tribosupplementation; tissue adhesion inhibition;
XX friction coefficient reduction; gene therapy; antiarthritic;
XX osteopathic; reverse transcription-PCR; RT-PCR primer; ss.
XX
XX Homo sapiens.
XX
XX WO200064930-A2.
XX
XX 02-NOV-2000.
XX

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PF 24-APR-2000; 2000WO-US10953.
XX
XX
PR 23-APR-1999; 99US-0298970.
XX
XX (RHOD-) RHODE ISLAND HOSPITAL LIFESPAN PARTNER.
XX
XX Jay GD;
XX
XX WPI; 2001-024673/03.
XX
XX Novel tribonectin polypeptide useful as lubricant for treating
PT osteoarthritis, comprises O-linked lubricating moiety -
XX
XX Example 1; Page 23; 47pp; English.
XX
XX The invention relates to a human tribonectin which is a product of
CC alternative splicing of the human MSF (megakaryocyte stimulating factor)
CC gene. The tribonectin has at least one O-linked oligosaccharide
CC lubricating moiety and has a polypeptide sequence comprising 1-76
CC repeats of a motif having at least 50% identity to the sequence KEPAPTT
CC (AA829774). The invention also relates to a nucleic acid encoding a
CC human MSF-derived tribonectin; a biocompatible composition comprising a
CC human tribonectin for inhibiting tissue adhesion formation; and a method
CC of diagnosing osteoarthritis or a predisposition to osteoarthritis by
CC measuring the amount of MSF or its fragment in a biological sample of a
CC mammal, wherein an increased amount of MSF compared to a control
CC indicates the presence of or predisposition to developing
CC osteoarthritis. The tribonectin and DNA encoding it are useful in the
CC treatment of osteoarthritis, where they may be used for lubricating
CC mammalian joints, such as articulating joints of humans, dogs or horses.
CC The tribonectin, when formulated as a membrane, foam, gel or fibre, is
CC useful for inhibiting adhesion between two surfaces such as the injured
CC tissues of a mammal, where the injury is caused by a surgical insertion
CC of trauma, or an artificial device e.g., an orthopaedic implant. In
CC particular, one of the surfaces is pericardial tissue. DNA encoding a
CC tribonectin may be used in gene therapy. The present sequence represents
CC a human MSF gene reverse transcription-PCR (RT-PCR) primer used in an
CC exemplification of the invention.
XX
XX Sequence 28 BP; 5 A; 7 C; 6 G; 10 T; 0 other;
SQ
Query Match 100.0%; Score 10; DB 22; Length 28;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db |||||
12 CTTCTCTTTT 21
RESULT 97
AAF05757/c
ID AAF05757 standard; RNA; 29 BP.
XX
XX AAF05757;
XX
XX 16-FEB-2001 (first entry)
XX
XX Hammerhead ribozyme #2976.
XX
XX Ribozyme; erythropoietin; granulocyte colony stimulating factor;
XX interferon alpha; ss.
XX
XX Homo sapiens.
XX
XX WO200061729-A2.
XX
XX 19-OCT-2000.
XX
XX 11-APR-2000; 2000WO-US09721.
XX
XX 12-APR-1999; 99US-0129390.
XX
XX

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PA (RIBO-) RIBOZYME PHARM INC.
XX
XX Blatt L, Zwick M, Pavco P, McSwiggen J;
XX
XX WPI; 2000-647423/62.
XX
XX Enzymatic and antisense nucleic acid inhibition of repressor genes,
PT useful for producing e.g. granulocyte colony stimulating factor
PT protein, interferon alpha and erythropoietin -
XX
XX Claim 48; Page 124; 164pp; English.
XX
XX The present invention relates to enzymatic and antisense nucleic acid
CC molecules that act as inhibitors of the expression of repressor genes
CC encoding the TR2 Orphan receptor, EAR3/COUP-IF-1, the GATA
CC transcription factor gene, IRF-2 and/or the CAAT Displacement
CC Protein (CDP). Inhibition of the repressors removes prevents
CC inhibition (and consequently increases expression of) genes involved in
CC the production of erythropoietin, granulocyte colony stimulating factor
CC protein and interferon alpha.
XX
XX Sequence 29 BP; 10 A; 4 C; 9 G; 5 U; 1 other;
SQ
Query Match 100.0%; Score 10; DB 21; Length 29;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db |||||
29 CTTCTCTTTT 20
RESULT 98
AAA40663
ID AAA40663 standard; DNA; 29 BP.
XX
XX AAA40663;
XX
XX 15-AUG-2000 (first entry)
XX
XX Human CD36 polymorphism sequence variant oligonucleotide SEQ ID NO:133.
XX
XX Human; rat; CD36; SHR; spontaneous hypertensive rat; diagnosis;
XX therapy; screening; polymorphism; variant; detection; mutant;
XX blood; mutation; insulin; glucose metabolism; fatty acid metabolism;
XX catecholamine; malaria; infection; parasite; antiparasitic;
XX antidiabetic; primer; ss.
XX
XX Homo sapiens.
XX
XX Synthetic.
XX
XX WO200019883-A2.
XX
XX 13-APR-2000.
XX
XX 07-OCT-1999; 99WO-US23418.
XX
XX 07-OCT-1998; 98US-0167750.
XX
XX 28-DEC-1998; 98US-0221222.
XX
XX 17-MAR-1999; 99US-0270542.
XX
XX (MEDI-) MEDICAL RES COUNCIL.
XX (SCIO-) SCIOS INC.
XX (AITM/) AITMAN T J.
XX (SCOT/) SCOTT J.
XX (STAN/) STANTON L W.
XX
XX Aitman TJ, Scott J, Stanton LW;
XX
XX WPI; 2000-303596/26.
XX
XX Nucleic acids encoding mutant CD36 proteins useful for preventing,
PT diagnosing and treating parasitic infections, especially malaria -
PT

```


XX PS Claim 26; Page 90; 167pp; English.

XX CC The present invention describes isolated nucleic acid molecules (A) encoding mutant CD36 proteins (B). Parasites such as Plasmodium falciparum (the major cause of malaria) are unable to utilise the mutated proteins to gain entry to, and infect cells. The mutant CD36 proteins do not function correctly preventing parasites utilising them to infect cells. The nucleic acids may be used for the recombinant production of mutant CD36 proteins according to standard methodologies. They may be used in this way to prevent and treat parasitic infections. CC That utilise the CD36 protein to infect cells, such as P. falciparum, CC the major cause of malaria. For example, the protein may be used to CC identify modulators of CD36 expression and activity or a patient's CD36 CC DNA may be screened to determine whether there are any mutations present CC that may confer resistance to parasitic infections. The proteins and CC nucleic acids may also be used to prevent, diagnose and treat diseases CC associated with defects in insulin action and/or glucose metabolism CC and/or fatty acid metabolism and/or catecholamine action in subjects CC possessing mutations in the CD36 genes. AAA40606 to AAA40759, and CC AAB02515 to AAB02564, represent nucleotide and amino acid sequences CC respectively which are used in the exemplification of the present CC invention.

XX SQ Sequence 29 BP; 1 A; 5 C; 10 G; 13 T; 0 other;

Query Match 100.0%; Score 10; DB 21; Length 29;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17

RESULT 99
AAA40662
ID AAA40662 standard; DNA; 30 BP.
XX AC AAA40662;
XX DT 15-AUG-2000 (first entry)
XX DE Human CD36 polymorphism sequence variant oligonucleotide SEQ ID NO:132.
XX KW Human; rat; CD36; SHR; spontaneous hypertensive rat; diagnosis;
KW therapy; screening; polymorphism; variant; detection; mutant;
KW blood; mutation; insulin; glucose metabolism; fatty acid metabolism;
KW catecholamine; malaria; infection; parasite; antiparasitic;
KW antidiabetic; primer; ss.
XX OS Homo sapiens.
OS Synthetic.
XX PN WO200019883-A2.
XX PD 13-APR-2000.
XX PF 07-OCT-1999; 99WO-US23418.
XX PR 07-OCT-1998; 98US-0167750.
PR 28-DEC-1998; 98US-0221222.
PR 17-MAR-1999; 99US-0270542.
XX (MEDI-) MEDICAL RES COUNCIL.
PA (SCIO-) SCIOS INC.
PA (AITM/) AITMAN T J.
PA (SCOT/) SCOTT J.
PA (STAN/) STANTON L W.
XX Aitman TJ, Scott J, Stanton LW;
XX WPI; 2000-303596/26.

XX PT Nucleic acids encoding mutant CD36 proteins useful for preventing, diagnosing and treating parasitic infections, especially malaria -

XX PS Claim 26; Page 90; 167pp; English.

XX CC The present invention describes isolated nucleic acid molecules (A) encoding mutant CD36 proteins (B). Parasites such as Plasmodium falciparum (the major cause of malaria) are unable to utilise the mutated proteins to gain entry to, and infect cells. The mutant CD36 proteins do not function correctly preventing parasites utilising them to infect cells. The nucleic acids may be used for the recombinant production of mutant CD36 proteins according to standard methodologies. They may be used in this way to prevent and treat parasitic infections. CC That utilise the CD36 protein to infect cells, such as P. falciparum, CC the major cause of malaria. For example, the protein may be used to CC identify modulators of CD36 expression and activity or a patient's CD36 CC DNA may be screened to determine whether there are any mutations present CC that may confer resistance to parasitic infections. The proteins and CC nucleic acids may also be used to prevent, diagnose and treat diseases CC associated with defects in insulin action and/or glucose metabolism CC and/or fatty acid metabolism and/or catecholamine action in subjects CC possessing mutations in the CD36 genes. AAA40606 to AAA40759, and CC AAB02515 to AAB02564, represent nucleotide and amino acid sequences CC respectively which are used in the exemplification of the present CC invention.

XX SQ Sequence 30 BP; 1 A; 5 C; 10 G; 14 T; 0 other;

Query Match 100.0%; Score 10; DB 21; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17

RESULT 100
AAA40664
ID AAA40664 standard; DNA; 30 BP.
XX AC AAA40664;
XX DT 15-AUG-2000 (first entry)
XX DE Human CD36 polymorphism sequence variant oligonucleotide SEQ ID NO:134.
XX KW Human; rat; CD36; SHR; spontaneous hypertensive rat; diagnosis;
KW therapy; screening; polymorphism; variant; detection; mutant;
KW blood; mutation; insulin; glucose metabolism; fatty acid metabolism;
KW catecholamine; malaria; infection; parasite; antiparasitic;
KW antidiabetic; primer; ss.
XX OS Homo sapiens.
OS Synthetic.
XX PN WO200019883-A2.
XX PD 13-APR-2000.
XX PF 07-OCT-1999; 99WO-US23418.
XX PR 07-OCT-1998; 98US-0167750.
PR 28-DEC-1998; 98US-0221222.
PR 17-MAR-1999; 99US-0270542.
XX (MEDI-) MEDICAL RES COUNCIL.
PA (SCIO-) SCIOS INC.
PA (AITM/) AITMAN T J.
PA (SCOT/) SCOTT J.
PA (STAN/) STANTON L W.
XX Aitman TJ, Scott J, Stanton L W.

PI Aitman TJ, Scott J, Stanton LW;
XX
DR WPI; 2000-303596/26.
XX
XX Nucleic acids encoding mutant CD36 proteins useful for preventing,
PT diagnosing and treating parasitic infections, especially malaria -
PT
XX Claim 26; Page 90; 167pp; English.
PS
XX The present invention describes isolated nucleic acid molecules (A)
CC encoding mutant CD36 proteins (B). Parasites such as Plasmodium
CC falciparum (the major cause of malaria) are unable to utilise the
CC mutated proteins to gain entry to, and infect cells. The mutant CD36
CC proteins do not function correctly preventing parasites utilising them
CC to infect cells. The nucleic acids may be used for the recombinant
CC production of mutant CD36 proteins according to standard methodologies.
CC They may be used in this way to prevent and treat parasitic infections.
CC that utilise the CD36 protein to infect cells, such as P. falciparum,
CC the major cause of malaria. For example, the protein may be used to
CC identify modulators of CD36 expression and activity or a patient's CD36
CC DNA may be screened to determine whether there are any mutations present
CC that may confer resistance to parasitic infections. The proteins and
CC nucleic acids may also be used to prevent, diagnose and treat diseases
CC associated with defects in insulin action and/or glucose metabolism
CC and/or fatty acid metabolism and/or catecholamine action in subjects
CC possessing mutations in the CD36 genes. AAA40606 to AAA40759, and
CC AAB02515 to AAB02564, represent nucleotide and amino acid sequences
CC respectively which are used in the exemplification of the present
CC invention.
XX
SQ Sequence 30 BP; 1 A; 5 C; 10 G; 14 T; 0 other;

Query Match 100.0%; Score 10; DB 21; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.7e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17

Search completed: October 28, 2003, 17:17:58
Job time : 287 secs

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OM nucleic - nucleic search, using sw model

Run on: October 28, 2003, 15:14:10 ; Search time 2088 Seconds
(without alignments)
116.401 Million cell updates/sec

Title: US-09-335-032-71
Perfect score: 10
Sequence: 1 cttctctttt 10

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 22781392 seqs, 12152238056 residues

Word size : 0

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 500 summaries

Database :

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4: em_estmu:*
5: em_estov:*
6: em_estpl:*
7: em_estro:*
8: em_hcc:*
9: gb_est1:*
10: gb_est2:*
11: gb_hcc:*
12: gb_est3:*
13: gb_est4:*
14: gb_est5:*
15: em_estfun:*
16: em_estom:*
17: em_gss_hum:*
18: em_gss_inv:*
19: em_gss_pln:*
20: em_gss_vrt:*
21: em_gss_fun:*
22: em_gss_mam:*
23: em_gss_mus:*
24: em_gss_pro:*
25: em_gss_rod:*
26: em_gss_phg:*
27: em_gss_vrl:*
28: gb_gss1:*
29: gb_gss2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	10	100.0	25	29	CC457116 SALK 1066
2	10	100.0	28	9	AI686998 tp81e01.x
3	10	100.0	28	29	TA130812P
C 4	10	100.0	29	13	BQ590098

C	5	10	100.0	32	12	BI081179
C	6	10	100.0	32	29	AL941390
C	7	10	100.0	33	29	BZ763249
C	8	10	100.0	34	9	AU256102
C	9	10	100.0	37	28	AZ429862
C	10	10	100.0	37	28	AZ514585
C	11	10	100.0	39	28	AZ613373
C	12	10	100.0	40	9	AA916182
C	13	10	100.0	41	28	AZ375959
C	14	10	100.0	42	29	BI20830
C	15	10	100.0	42	29	BI24544
C	16	10	100.0	45	29	CC050354
C	17	10	100.0	45	29	BI20970
C	18	10	100.0	49	9	AA922976
C	19	10	100.0	49	9	AI564984
C	20	10	100.0	49	29	BZ357069
C	21	10	100.0	50	9	AU102691
C	22	10	100.0	51	29	BZ770334
C	23	10	100.0	51	29	AB082596
C	24	10	100.0	52	13	BQ667496
C	25	10	100.0	52	28	AZ327146
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C	31	10	100.0	58	9	AI343303
C	32	10	100.0	58	13	BQ595228
C	33	10	100.0	59	9	AU077187
C	34	10	100.0	59	28	AZ774339
C	35	10	100.0	59	29	BZ354535
C	36	10	100.0	59	29	AL762585
C	37	10	100.0	63	9	AU258677
C	38	10	100.0	64	10	AW874904
C	39	10	100.0	64	10	AW874934
C	40	10	100.0	64	10	BE239248
C	41	10	100.0	64	10	BE239284
C	42	10	100.0	64	10	BE239293
C	43	10	100.0	64	10	BE636299
C	44	10	100.0	64	10	BE636308
C	45	10	100.0	64	10	BE636320
C	46	10	100.0	64	10	BE636344
C	47	10	100.0	64	10	BE636363
C	48	10	100.0	64	10	BE636369
C	49	10	100.0	64	10	BE636392
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C	51	10	100.0	64	10	BE636451
C	52	10	100.0	64	10	BE636456
C	53	10	100.0	64	10	BE636464
C	54	10	100.0	64	10	BE636472
C	55	10	100.0	64	10	BF118488
C	56	10	100.0	64	12	BI097426
C	57	10	100.0	64	12	BI142401
C	58	10	100.0	64	12	BI142408
C	59	10	100.0	64	12	BI142422
C	60	10	100.0	64	12	BI142452
C	61	10	100.0	64	28	AZ821452
C	62	10	100.0	65	14	CB030041
C	63	10	100.0	65	28	AZ514453
C	64	10	100.0	65	29	CC179177
C	65	10	100.0	65	29	AG216203
C	66	10	100.0	65	29	AL769814
C	67	10	100.0	65	29	AL949773
C	68	10	100.0	65	29	TA129A12P
C	69	10	100.0	66	14	W85242
C	70	10	100.0	66	28	AZ514401
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C	73	10	100.0	68	12	BM447291
C	74	10	100.0	68	29	AL757142
C	75	10	100.0	70	9	AL651716
C	76	10	100.0	70	9	AA255635
C	77	10	100.0	70	14	H45651

BI081179	602879191
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BZ763249	SALK 1157
AU256102	AU256102
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AZ514585	IM0361B16
AZ613373	IM0441M12
AA916182	OG34B06.s
AZ375959	IM0129A12
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BI24544	Danio rer
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BI20970	Danio rer
AA922976	ok77e06.s
AI564984	tg53b02.x
BZ357069	SALK_1301
AU102691	AU102691
BZ770334	SALK 1432
AB082596	Drosophil
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AZ327146	IM0050015
BE569086	601339390
BH908756	SALK_0504
BH864698	SALK_0967
BI322307	Kx19G11.Y
CB917098	VVD118G10
AI343303	tb92G09.X
BQ595228	E012710-0
AU077187	AU077187
AZ774339	2M0003005
BZ354535	SALK 1252
AL762585	Arabidops
AU258677	AU258677
AW874904	SWYACAL04
AW874934	SWYACAL04
BE239248	SMOVL2CAS
BE239284	SMOVL2CAS
BE239293	SMOVL2CAS
BE636299	SMOVL2CAS
BE636308	SMOVL2CAS
BE636320	SMOVL2CAS
BE636344	SMOVL2CAS
BE636363	SMOVL2CAS
BE636369	SMOVL2CAS
BE636392	SMOVL2CAS
BE636430	SMOVL2CAS
BE636451	SMOVL2CAS
BE636456	SMOVL2CAS
BE636464	SMOVL2CAS
BE636472	SMOVL2CAS
BF118488	SMOVL3CAN
BI097426	SMOV3MCAM
BI142401	SMOV3MCAM
BI142408	SMOV3MCAM
BI142422	SMOV3MCAM
BI142452	SMOV3MCAM
AZ821452	2M0094B14
CB030041	TG8STYd1
AZ514453	IM0361N14
CC179177	SALK 0591
AG216203	Drosophil
AL769814	Arabidops
AL949773	Arabidops
AL463978	T. brucei
W85242	mf52h08.r1
AZ514401	IM0361K04
AI442885	sa28b08.X
CD029241	mgns007XA
BM447291	DSA008D10
AL757142	Arabidops
AL651716	AL651716
AA255635	zs31f07.r
H45651	yn97d02.sl

C 78	10	100.0	70	28	AZ918371	1006004B0	C 151	10	100.0	94	29	CC458005	CC458005	SALK_1147
C 79	10	100.0	71	14	D78209	D78209 EST	C 152	10	100.0	95	9	A1955060	A1955060	wq60c05.x
C 80	10	100.0	71	29	BZ770025	SALK_1429	C 153	10	100.0	95	9	AW713887	AW713887	h3jg02ne.f
C 81	10	100.0	72	9	AA184862	mu51b11.r	C 154	10	100.0	95	29	DR24C7S	DR24C7S	DA10 rer
C 82	10	100.0	72	29	BX285944	Arabidops	C 155	10	100.0	96	13	BUR63237	BUR63237	S025F09 P
C 83	10	100.0	72	9	AA262253	zr70g04.r	C 156	10	100.0	96	28	AF179181	AF179181	AF179181
C 84	10	100.0	73	9	AA976531	SWYD25CAU	C 157	10	100.0	96	29	BZ288800	SALK_0221	BZ288800
C 85	10	100.0	73	9	AW600116	SWL4CAK10	C 158	10	100.0	97	9	A1255137	qv46d01.x	A1255137
C 86	10	100.0	73	9	AW626514	SWOVL3CAN	C 159	10	100.0	97	9	A1360659	qx59d01.x	A1360659
C 87	10	100.0	73	9	AW626555	SWOVL3CAN	C 160	10	100.0	97	28	AZ461670	1M0267F11	AZ461670
C 88	10	100.0	73	9	AW651817	SWYD25CAU	C 161	10	100.0	97	29	BZ383278	SALK_1323	BZ383278
C 89	10	100.0	73	10	BG310475	SWOV3MCAU	C 162	10	100.0	97	29	CC029290	3591_1.10	CC029290
C 90	10	100.0	73	10	AW874933	SWYACAL04	C 163	10	100.0	97	29	AG229920	LOTus_Tad	AG229920
C 91	10	100.0	73	10	BE420470	SWOVL2CAS	C 164	10	100.0	98	10	BE621613	601493758	BE621613
C 92	10	100.0	73	10	BE420471	SWOVL2CAS	C 165	10	100.0	98	28	BH862099	SALK_0887	BH862099
C 93	10	100.0	73	10	BE420480	SWOVL2CAS	C 166	10	100.0	98	29	AL1770908	Arabidops	AL1770908
C 94	10	100.0	73	10	BE638405	SWOVL2CAS	C 167	10	100.0	99	9	A1110549	SWOVL3CAN	A1110549
C 95	10	100.0	73	10	BF228818	SWOVL3CAN	C 168	10	100.0	99	9	AT006315	AT006315	AT006315
C 96	10	100.0	73	14	CB886667	CB886667	C 169	10	100.0	99	28	AZ333242	1M0062D14	AZ333242
C 97	10	100.0	74	29	BZ660902	SALK_0243	C 170	10	100.0	99	29	CC459096	SALK_1247	CC459096
C 98	10	100.0	75	14	CA849383	K111903.y	C 171	10	100.0	100	9	AI197630	ue45d06.r	AI197630
C 99	10	100.0	75	29	TA77C02P	AL460777 T. brucei	C 172	10	100.0	100	9	AW231456	687061G10	AW231456
C 100	10	100.0	76	9	AA615345	vo61d12.r	C 173	10	100.0	100	10	BG153513	naq49f05.	BG153513
C 101	10	100.0	76	10	BF055674	7i68d03.y	C 174	10	100.0	100	10	BE067879	RC0-BT036	BE067879
C 102	10	100.0	76	28	AZ309821	1M0017C11	C 175	10	100.0	100	10	BE153449	PM2-HT033	BE153449
C 103	10	100.0	76	29	AL768539	Arabidops	C 176	10	100.0	100	13	BU097798	946119D08	BU097798
C 104	10	100.0	76	29	CNS02MOR	AL204372 Tetraodon	C 177	10	100.0	100	14	CB170638	CKV602600	CB170638
C 105	10	100.0	77	13	BQ456810	ke31c12.y	C 178	10	100.0	100	14	D82659	HUMHBC3277	D82659
C 106	10	100.0	77	14	CB099465	ks10h09.y	C 179	10	100.0	101	9	AL871534	AL871534	AL871534
C 107	10	100.0	77	14	CB277589	ks37g04.y	C 180	10	100.0	101	9	AA236344	zr51f06.r	AA236344
C 108	10	100.0	77	14	CB277809	ks39g02.y	C 181	10	100.0	101	10	BE500183	WHE0980.F	BE500183
C 109	10	100.0	77	29	AL758789	Arabidops	C 182	10	100.0	102	10	BF354509	CM1-HT076	BF354509
C 110	10	100.0	78	29	AL762583	AL762583 Arabidops	C 183	10	100.0	102	12	BN090933	1g17h06.x	BN090933
C 111	10	100.0	80	9	AA571271	AA571271 v190f03.r	C 184	10	100.0	102	13	BU893309	P076A10.P	BU893309
C 112	10	100.0	80	29	BZ289251	BZ289251 SALK_0226	C 185	10	100.0	102	28	BH812258	BOGMS84TR	BH812258
C 113	10	100.0	81	28	BH855764	SALK_0846	C 186	10	100.0	102	28	BH812258	SALK_0615	BH812258
C 114	10	100.0	82	28	BH911702	BH911702 SALK_0716	C 187	10	100.0	103	9	AW361423	RC3-CT025	AW361423
C 115	10	100.0	82	29	AL765739	AL765739 Arabidops	C 188	10	100.0	103	10	AW888322	M43_Rac.d	AW888322
C 116	10	100.0	83	10	BF507234	4682P-15a	C 189	10	100.0	103	12	BI127881	G067F38Y	BI127881
C 117	10	100.0	83	14	F14596	F14596 SSO4D05 Por	C 190	10	100.0	103	12	BJ327682	BJ327682	BJ327682
C 118	10	100.0	83	14	F37812	F37812 HSPD06839 H	C 191	10	100.0	103	14	CA812476	CA48LN061	CA812476
C 119	10	100.0	83	29	AL941379	Arabidops	C 192	10	100.0	103	28	BH700682	BOMMG89TF	BH700682
C 120	10	100.0	84	14	CB001243	VVB002H02	C 193	10	100.0	103	28	BH818088	BACPL0-F	BH818088
C 121	10	100.0	84	28	BH251470	SALK_0116	C 194	10	100.0	103	29	BZ314704	hzz2e09.b	BZ314704
C 122	10	100.0	84	28	BH809934	SALK_0367	C 195	10	100.0	104	9	AI407102	EST235390	AI407102
C 123	10	100.0	84	29	CNS04INL	AL270426 Tetraodon	C 196	10	100.0	104	9	AA170839	JTH131 HT	AA170839
C 124	10	100.0	85	12	BI703760	ks18g12.y	C 197	10	100.0	104	9	AW062642	RC0-CT009	AW062642
C 125	10	100.0	85	13	BQ252968	sao04d04.	C 198	10	100.0	104	10	BF747371	RC3-BT033	BF747371
C 126	10	100.0	85	14	U44279	ENU44279 As	C 199	10	100.0	104	10	BF833447	PM1-HT092	BF833447
C 127	10	100.0	85	28	AZ855489	AZ855489 2M0159L04	C 200	10	100.0	104	10	BF991035	CM1-GN016	BF991035
C 128	10	100.0	85	28	BH862052	SALK_0887	C 201	10	100.0	104	12	BI014338	CM4-ET014	BI014338
C 129	10	100.0	85	29	EX288399	EX288399 Arabidops	C 202	10	100.0	104	12	BI044504	PM3-OT020	BI044504
C 130	10	100.0	86	9	AA067973	mm25a10.r	C 203	10	100.0	104	12	BI179001	EST519946	BI179001
C 131	10	100.0	86	9	AJ547920	AJ547920	C 204	10	100.0	104	12	BJ390565	BJ390565	BJ390565
C 132	10	100.0	86	10	BG673194	BG673194 DRNBD05	C 205	10	100.0	104	12	BM403561	zam4875.Z	BM403561
C 133	10	100.0	86	29	BI175324	BI175324 Danilo rer	C 206	10	100.0	104	13	BQ358421	CM0-HT091	BQ358421
C 134	10	100.0	89	9	AV567495	AV567495	C 207	10	100.0	104	13	BQ864530	SO4F12.F	BQ864530
C 135	10	100.0	89	28	BH251474	SALK_0116	C 208	10	100.0	104	29	CC027328	3591_1.59	CC027328
C 136	10	100.0	89	29	BZ292893	BZ292893 SALK_1285	C 209	10	100.0	105	9	AA068215	mm48g02.r	AA068215
C 137	10	100.0	90	13	BQ098870	BQ098870 ph23f03.y	C 210	10	100.0	105	9	AB036715	AB036715	AB036715
C 138	10	100.0	90	13	BQ582926	BQ582926 S015372-0	C 211	10	100.0	105	9	AA075027	zm83h05.s	AA075027
C 139	10	100.0	91	9	AA627648	ng51d08.s	C 212	10	100.0	105	9	AT006282	AT006282	AT006282
C 140	10	100.0	91	28	BH908214	BH908214 SALK_0484	C 213	10	100.0	105	9	AA485070	aa40b05.s	AA485070
C 141	10	100.0	92	9	AA585277	PTH020 HT	C 214	10	100.0	105	13	BQ325343	CM1-CI000	BQ325343
C 142	10	100.0	92	14	C20817	C20817 HUMGS000486	C 215	10	100.0	105	14	CA798394	Cac BL 72	CA798394
C 143	10	100.0	92	28	BH215979	BH215979 1006039C1	C 216	10	100.0	105	28	AZ797219	2M053015	AZ797219
C 144	10	100.0	92	29	CC179017	CC179017 SALK_0571	C 217	10	100.0	106	9	AI230637	EST227332	AI230637
C 145	10	100.0	93	28	BH864683	BH864683 SALK_0966	C 218	10	100.0	106	9	AU260000	AU260000	AU260000
C 146	10	100.0	93	28	BH864684	BH864684 SALK_0967	C 219	10	100.0	106	9	AV902900	AV902900	AV902900
C 147	10	100.0	93	28	BH864686	BH864686 SALK_0967	C 220	10	100.0	106	10	BE840762	MR2-SN000	BE840762
C 148	10	100.0	93	28	BH864689	BH864689 SALK_0967	C 221	10	100.0	106	12	BI322353	kk20a02.y	BI322353
C 149	10	100.0	94	12	BI174502	OSTF046C1	C 222	10	100.0	106	14	CD026264	NXSI_133	CD026264
C 150	10	100.0	94	28	BH851869	BH851869 SALK_0736	C 223	10	100.0	107	9	AI200462	qf93d09.x	AI200462

224	10	100.0	107	12	BJ358573	BJ358573	297	10	100.0	115	10	BF475908	BF475908
225	10	100.0	107	12	BJ389477	BJ389477	c 298	10	100.0	115	10	BG133296	BG133296
226	10	100.0	107	14	H23901	Ym75a05.r1	c 299	10	100.0	115	10	AW912033	AW912033
227	10	100.0	107	28	AZ565656	z11Ppf04	c 300	10	100.0	115	12	BM005397	BM005397
228	10	100.0	108	9	AW813619	RC3-ST019	c 301	10	100.0	115	12	BM026664	BM026664
229	10	100.0	108	14	T83754	Yd67h04.s1	c 302	10	100.0	115	28	AQ357469	AQ357469
230	10	100.0	108	28	BH118434	RPCI-24-2	c 303	10	100.0	116	9	AI252189	AI252189
231	10	100.0	108	28	BH416075	BH416075	c 304	10	100.0	116	9	AW394059	AW394059
232	10	100.0	109	9	AW753478	PM0-CT026	c 305	10	100.0	116	10	BG085551	BG085551
233	10	100.0	109	10	BE008587	RC5-BN015	c 306	10	100.0	116	28	BH078074	BH078074
234	10	100.0	109	10	BE012104	RC5-BN105	c 307	10	100.0	116	28	BH078074	BH078074
235	10	100.0	109	10	BE315587	NFO23E10L	c 308	10	100.0	117	13	BH080936	BH080936
236	10	100.0	109	13	BQ591622	E012616-0	c 309	10	100.0	117	28	BH105487	BH105487
237	10	100.0	109	14	CB405014	OSTR036B2	c 310	10	100.0	117	28	BH105487	BH105487
238	10	100.0	109	14	T91479	Yel8h09.r1	c 311	10	100.0	117	29	BZ377590	BZ377590
239	10	100.0	109	28	AQ080743	CIT-HSP-2	c 312	10	100.0	117	29	BZ764494	BZ764494
240	10	100.0	109	28	BH414927	1007040E0	c 313	10	100.0	118	9	AA934372	AA934372
241	10	100.0	110	9	AA762509	Vv84B12.r	c 314	10	100.0	118	9	AW001605	AW001605
242	10	100.0	110	9	AV903734	AV903734	c 315	10	100.0	118	9	AW001605	AW001605
243	10	100.0	110	10	BG310551	SNOW3MCAM	c 316	10	100.0	118	10	BF772244	BF772244
244	10	100.0	110	14	CB006382	VVC033B04	c 317	10	100.0	118	10	AW845662	AW845662
245	10	100.0	110	28	BH347115	CH230-120	c 318	10	100.0	118	10	BE998209	BE998209
246	10	100.0	110	28	BH637436	1008016E0	c 319	10	100.0	118	12	BI400446	BI400446
247	10	100.0	110	28	BH809571	KG07049-3	c 320	10	100.0	118	14	CA881406	CA881406
248	10	100.0	110	28	AQ385208	RPCI11-15	c 321	10	100.0	118	14	D82788	D82788
249	10	100.0	110	29	CC365311	PUCQP53TD	c 322	10	100.0	118	14	T48514	T48514
250	10	100.0	111	9	AL468239	Vv84B12.Y	c 323	10	100.0	118	28	AZ247438	AZ247438
251	10	100.0	111	9	AV427864	AV427864	c 324	10	100.0	118	28	AZ483588	AZ483588
252	10	100.0	111	12	BM092640	sah16g11.	c 325	10	100.0	118	28	BH340928	BH340928
253	10	100.0	111	13	BM109777	BM109777	c 326	10	100.0	118	29	CNS0120E	CNS0120E
254	10	100.0	111	14	CB385906	OSTF036B2	c 327	10	100.0	119	9	AV559784	AV559784
255	10	100.0	111	28	BZ115526	CH230-422	c 328	10	100.0	119	9	BF836144	BF836144
256	10	100.0	111	28	AQ482662	RPCI-11-2	c 329	10	100.0	119	10	BG338662	BG338662
257	10	100.0	111	29	BZ290112	SALK_0235	c 330	10	100.0	119	12	BG338662	BG338662
258	10	100.0	111	29	CC106912	CC106912	c 331	10	100.0	119	12	BM139880	BM139880
259	10	100.0	112	9	AA212505	mu79g03.r	c 332	10	100.0	119	14	CA847224	CA847224
260	10	100.0	112	9	AW384207	RC0-HT037	c 333	10	100.0	119	14	CB885142	CB885142
261	10	100.0	112	9	AA564935	nj24d11.s	c 334	10	100.0	119	28	BH542039	BH542039
262	10	100.0	112	9	BG604174	EST456372	c 335	10	100.0	119	29	AL754823	AL754823
263	10	100.0	112	10	BQ35562	SK4-0062	c 336	10	100.0	120	9	AA082593	AA082593
264	10	100.0	112	14	R47206	CB8-418 Sub	c 337	10	100.0	120	9	AA082593	AA082593
265	10	100.0	112	28	AZ904406	RPCI-24-2	c 338	10	100.0	120	9	AW393738	AW393738
266	10	100.0	112	28	AZ905862	RPCI-24-1	c 339	10	100.0	120	13	BQ312351	BQ312351
267	10	100.0	112	28	BH202354	SNL-58A5.	c 340	10	100.0	120	13	BU092901	BU092901
268	10	100.0	112	29	BZ734963	OGEDS70TC	c 341	10	100.0	120	13	BU092901	BU092901
269	10	100.0	112	29	CC058942	1117d03.9	c 342	10	100.0	120	14	AZ736188	AZ736188
270	10	100.0	112	29	AL946882	ArabiDops	c 343	10	100.0	120	28	BH195988	BH195988
271	10	100.0	113	9	AV970176	AV970176	c 344	10	100.0	120	28	BH195988	BH195988
272	10	100.0	113	9	AW384162	RC0-HT037	c 345	10	100.0	121	9	AA603854	AA603854
273	10	100.0	113	9	AW384195	RC0-HT037	c 346	10	100.0	121	10	BE197817	BE197817
274	10	100.0	113	10	BF735099	CM1-AN008	c 347	10	100.0	121	28	AZ086687	AZ086687
275	10	100.0	113	10	BE008812	CM4-BN016	c 348	10	100.0	121	28	AZ879966	AZ879966
276	10	100.0	113	12	BJ364111	BJ364111	c 349	10	100.0	121	28	AZ911833	AZ911833
277	10	100.0	113	12	BJ369833	BJ369833	c 350	10	100.0	121	28	AZ911833	AZ911833
278	10	100.0	113	12	BJ409035	BJ409035	c 351	10	100.0	121	29	BZ958145	BZ958145
279	10	100.0	113	13	BQ114930	EST600506	c 352	10	100.0	122	9	AL138752	AL138752
280	10	100.0	113	14	CA794594	CAc BL_14	c 353	10	100.0	122	9	AW129644	AW129644
281	10	100.0	113	28	AZ564228	RPCI-23-2	c 354	10	100.0	122	10	BF735541	BF735541
282	10	100.0	113	28	BH851383	SALK_0729	c 355	10	100.0	122	10	BF998919	BF998919
283	10	100.0	113	29	BZ358624	SALK_1330	c 356	10	100.0	122	10	BG093701	BG093701
284	10	100.0	113	29	AL751707	ArabiDops	c 357	10	100.0	122	10	BG347903	BG347903
285	10	100.0	114	9	AL830940	wj80b07.x	c 358	10	100.0	122	12	BG838320	BG838320
286	10	100.0	114	9	AL965770	sc75h05.Y	c 359	10	100.0	122	12	BF742650	BF742650
287	10	100.0	114	9	AW384183	RC0-HT037	c 360	10	100.0	122	12	BJ363631	BJ363631
288	10	100.0	114	9	AW695613	NFO97A045	c 361	10	100.0	122	12	BU541349	BU541349
289	10	100.0	114	10	AW920975	EST522279	c 362	10	100.0	122	13	BU574463	BU574463
290	10	100.0	114	12	BJ358564	BJ358564	c 363	10	100.0	122	14	CB515109	CB515109
291	10	100.0	114	12	BM329382	PIC1_37_B	c 364	10	100.0	122	29	BX245707	BX245707
292	10	100.0	114	28	BH212127	SALK_0071	c 365	10	100.0	123	9	AA749641	AA749641
293	10	100.0	114	28	BH226094	1006130A0	c 366	10	100.0	123	9	AU222440	AU222440
294	10	100.0	114	28	BH827449	BACPP25-O	c 367	10	100.0	123	10	BF353080	BF353080
295	10	100.0	115	9	AT862639	wj27b08.x	c 368	10	100.0	123	10	BF355020	BF355020
296	10	100.0	115	9	AV984486	AV984486	c 369	10	100.0	123	12	BI433964	BI433964

C	370 c	371		10	100.0	123	BJ389997	BJ389997	CB458495	717151 MA	c	443	10	100.0	133	9	AI371623	AI371623 tb72g04.x
		123	14	CB458495	717151 MA							444	10	100.0	133	9	AI612429	AI612429 48608A04
		123	14	AZ079931	RPCI-23-4							445	10	100.0	133	9	AVI42453	AVI42453 AVI42453
		123	28	AZ091008	RPCI-23-4							446	10	100.0	133	10	BF835600	BF835600 RC4-HT089
		374	28	AZ289378	RPCI-23-4							447	10	100.0	133	10	BG135525	PQ3_0.219
		375	28	BH664559	BOMJU79TR							448	10	100.0	133	10	BF086053	CM3-CN004
		376	28	CC426132	PURKR33TB							449	10	100.0	133	10	BF249583	pah1hl.y
		377	29	EX289216	Arabidopsys							450	10	100.0	133	12	BG790316	sae67n03.
		378	9	ATW004196	ATW004196							451	10	100.0	133	12	EIO44208	PM4-COT20
		124	9	AW579502	MRO-HTO16							452	10	100.0	133	12	BM026751	fq87h12-y
		124	12	BE182022	CM1-HTO64							453	10	100.0	133	12	BM026872	ft10g02-y
		380	12	BJ325862	BJ325862							454	10	100.0	133	28	AZ367678	IM0117L02
		381	12	BJ325862	BJ325862							455	10	100.0	133	28	AZ367678	IM0117L02
		382	12	BJ416022	BJ416022							456	10	100.0	133	29	CC531659	PURKQ54TB
		383	12	CB914818	VVD09YTC04							457	-	100.0	134	9	AU210635	Oryza sat
		384	28	AZ233447	RPCI-23-9							458	10	100.0	134	9	AV165076	AV165076
		385	10	AZ345595	IM0080017							459	10	100.0	134	9	AV165076	AV165076
		386	28	AZ345595	IM0080017							460	10	100.0	134	9	AV165076	AV165076
		387	10	AL857650	wk9SNO9.X'							461	10	100.0	134	9	AM706651	sk01b03.y
		388	14	CA482726	LUP12010B							462	10	100.0	134	12	BM186774	fvT7f06.Y
		389	28	AQ987448	RPCI-23-3							463	10	100.0	134	12	BM537444	ha82d06.g
		390	28	AZ240596	RPCI-23-8							464	10	100.0	134	14	BY730099	BY730099
		391	28	BZ132001	CH230-345							465	10	100.0	134	14	CB274960	kut73e09.y
		392	9	AW436667	77088 MAR							466	10	100.0	134	14	CB274960	kut73e09.y
		393	10	BG724677	REESTed86							467	10	100.0	134	14	Z18441	ATT50816 Gr
		394	12	BJ388788	BJ388788							468	10	100.0	134	28	BH315791	HQ537522
		395	12	BQ367528	MRI-GN017							469	10	100.0	134	28	BH315791	HQ537522
		396	12	BM068489	BM068489							470	10	100.0	134	29	CE769428	SALK_1421
		397	10	CB931669	r161of.Y							471	10	100.0	134	29	CC119937	NDL_70G16
		398	28	AZ100181	RPCI-2													

ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids ; eurosids II; Brassicales; Brassicaceae; Arabidopsi.

REFERENCE
AUTHORS 1 (bases 1 to 25)
Alonso, J.M., Leisse, T.J., Batajag, P., Chen, H., Cheuk, R., Gadrinab , C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prednis, L., Shinn, P., Zimmerman, J. and Ecker, J.R.
TITLE A Sequence-Indexed Library of Insertion Mutations in the Arabidopsis Genome
JOURNAL Unpublished
COMMENT Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGNAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: eckersalk.edu
This is single pass sequence recovered from the left border of TDNA.
Class: TDNA tagged.
Location/Qualifiers
1. .25
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/db_xref="taxon:3702"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines each of which contains one or more TDNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at http://signal.salk.edu/tdna_protocols.html"

BASE COUNT 16 a 1 c 7 g 1 t
ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.5e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 22 CTTCTCTTTT 13

RESULT 2
AI686998
LOCUS AI686998 28 bp mRNA linear EST 14-DEC-1999
DEFINITION tp81e01.x1 NCI CGAP Ut3 Homo sapiens cDNA clone IMAGE:2205720 3' similar to TR:054875 054875 MYOTONIC DYSTROPHY KINASE-RELATED CDC42-BINDING KINASE MRCK-BETA. ; contains element MER17 repetitive element ; mRNA sequence.
ACCESSION AI686998.1 GI:4898292
VERSION AI686998.1
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
AUTHORS 1 (bases 1 to 28)
NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
JOURNAL Unpublished
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality
Insert Length: 2144 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 1.
Location/Qualifiers
1. .28
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone_lib="IMAGE:2205720"
/tissue_type="poorly-differentiated endometrial adenocarcinoma, 2 pooled tumors"
/lab_host="DH10B"
/clone_lib="NCI CGAP Ut3"
/note="Organ: uterus; Vector: pCMV-SPORT6; Site 1: SalI; Site 2: NotI; Cloned unidirectionally. Primer: Oligo dt. Average insert size 1.45 kb. Life Technologies catalog #:
11541-018"

BASE COUNT 4 a 12 c 0 g 12 t
ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 28;
Best Local Similarity 100.0%; Pred. No. 1.5e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 9 CTTCTCTTTT 18

RESULT 3
TA130B12P
LOCUS TA130B12P 28 bp DNA linear GSS 13-DEC-2000
DEFINITION T. brucei sheared genomic DNA clone 130b12, forward sequence, genomic survey sequence.
ACCESSION AL464095
VERSION AL464095.1 GI:11834358
KEYWORDS GSS.
SOURCE Trypanosoma brucei
ORGANISM Trypanosoma brucei
Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae; Trypanosoma.
REFERENCE
AUTHORS 1 (bases 1 to 28)
Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R., Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L., Melville, S.E., Rajadream, M.A. and Barrell, B.G.
Direct Submission
Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and nh@sanger.ac.uk
Constructed at the Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of Trypanosoma brucei (TREU927/4 GUTAT 10.1) was mechanically sheared to give a tight size distribution (4 kb). The v + i method used for the library construction is described in detail in Smith, H. and Venter, J.C. (Making small insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaudin and B. Barrell, Oxford University Press, 1999).
Email: nelsaved@tigr.org
Details of T. brucei sequencing at the Sanger Centre are available at http://www.sanger.ac.uk/Projects/T_brucei/.
Location/Qualifiers
1. .28
/organism="Trypanosoma brucei"
/mol_type="genomic DNA"
/db_xref="taxon:5691"

FEATURES
source

```

BASE COUNT      2 a      12 c      2 g      12 t
ORIGIN
Query Match      100.0%; Score 10; DB 29; Length 28;
Best Local Similarity 100.0%; Pred. No. 1.5e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 19 CTTCTCTTTT 28

RESULT 4
BQ590098/c
LOCUS
DEFINITION      BQ590098      29 bp      mRNA      linear      EST 06-DEC-2002
cDNA clone 024-019-019-T7 MP12-ADIS-024-storage root Beta vulgaris
ACCESSION      BQ590098
VERSION
KEYWORDS
SOURCE
ORGANISM
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Caryophyllales; Amaranthaceae; Beta.
REFERENCE
AUTHORS      Herwig,R., Schulz,B., Weisshaar,B., Hennig,S., Steinfath,M.,
Drungowski,M., Stahl,D., Wruck,W., Menze,A., O'Brien,J., Lehrach,H.
and Radelof,U.
TITLE
Construction of a 'unigene' cDNA clone set by oligonucleotide
fingerprinting allows access to 25 000 potential sugar beet genes
JOURNAL
COMMENT
Contact: Weisshaar B
ADIS DNA core facility at MP12
Max-Planck-Institute for Plant Breeding Research
Carl-von-Linne Weg 10, 50829 Koeln, Germany
Fax: 00492215062851
Email: weisshaar@mpiz-koeln.mpg.de
Insert Length: 29 Std Error: 0.00
Plate: 19 row: 0 column: 19
Seq primer: T7: GTAATACGACTCATTATAGGC.
FEATURES
source
1..29
Location/Qualifiers
/organism="Beta vulgaris"
/mol_type="mRNA"
/cultivar="KWS2320 (double haploid, monogerm breeding line
)"
/db_xref="GABI:190019"
/db_xref="taxon:161934"
/clone="024-019-019"
/tissue_type="storage root"
/lab_host="EMDH10B"
/clone_lib="MP12-ADIS-024-storage root"
/notes="Vector: pCMVSPORT6; Site 1: Sali; Site 2: NotI;
cDNA library from sugar beet, library provided by KWS
Kleinwanzlebener Saatucht AG Einbeck, Germany, contact:
b.schulz@kws.de; cloning sites Sali-NotI, primer sites and
orientation:
SP6-Sali-CCAGCGTCG-5prime-cDNA-polyA-CC-NotI-T7; Note:
Sequencing granted in the context of the GABI-Beet project
, local PI: Dr. Katharina Schneider, coordinator: Prof.
Christian Jung; Sequence submission managed by
RZPD/GABI-Primary database: http://gabi.rzpd.de"
BASE COUNT      15 a      0 c      14 g      0 t
ORIGIN
Query Match      100.0%; Score 10; DB 13; Length 29;
Best Local Similarity 100.0%; Pred. No. 1.5e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 19 CTTCTCTTTT 28

RESULT 5
BQ590098/c
LOCUS
DEFINITION      BQ590098      32 bp      mRNA      linear      EST 20-JUN-2001
cDNA clone 024-019-019-T7 MP12-ADIS-024-storage root Beta vulgaris
ACCESSION      BQ590098
VERSION
KEYWORDS
SOURCE
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Gilbert Smith, Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone Distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM11059 row: e column: 23
High quality sequence stop: 32.
FEATURES
source
1..32
Location/Qualifiers
/organism="Mus musculus"
/mol_type="mRNA"
/strain="FVB/N-3"
/db_xref="taxon:10090"
/clone="IMAGE:5010814"
/tissue_type="tumor, biopsy sample"
/dev_stage="5 months"
/lab_host="DH10B"
/clone_lib="NCI CGAP Mam2"
/notes="Organ: mammary; Vector: pCMV-SPORT6; Site 1: Sali;
Site 2: NotI; Cloned unidirectionally. Primer: Oligo dT.
Library constructed by Life Technologies. Investigator
providing samples: Gilbert Smith, NIH"
BASE COUNT      19 a      5 c      8 g      0 t
ORIGIN
Query Match      100.0%; Score 10; DB 12; Length 32;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 20 CTTCTCTTTT 11

RESULT 6
AL941390
LOCUS
DEFINITION      AL941390      32 bp      DNA      linear      GSS 24-OCT-2002
Arabidopsis thaliana T-DNA flanking sequence GK-257B04-014921,
genomic survey sequence.
ACCESSION      AL941390
VERSION
KEYWORDS
SOURCE
ORGANISM
Arabidopsis thaliana (thale cress)
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
REFERENCE
AUTHORS      Strizhov,N., Li,Y., Rosso,M., Viehoever,P., Dekker,K., Saedler,H.
and Weisshaar,B.

```


TITLE A pipeline for automated high-throughput generation of ESTs (flanking sequence tags) from Arabidopsis thaliana T-DNA transformed lines

JOURNAL Unpublished

REFERENCE 2

AUTHORS Rosso, M., Strizhov, N., Li, Y., Reiss, B., Dekker, K. and Weissshaar, B.

TITLE A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat) for flanking sequence tag based reverse genetics

JOURNAL Unpublished

REFERENCE 3 (bases 1 to 32)

AUTHORS Rosso, M., Li, Y., Strizhov, N. and Weissshaar, B.

TITLE Direct Submission

JOURNAL Submitted (21-OCT-2002) Weissshaar B., Max-Planck-Institut fuer Zuechtungsforschung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany

COMMENT This sequence is recovered from the left border of the T-DNA. It indicates an insertion close to or within gene At5g42900. The sequences are generated at the MPI for Plant Breeding Research in the context of the GABI-Kat project. GABI-Kat is part of the German Plant Genomics program designated 'GABI'. Information on line availability can be found at: <http://www.mpiz-koeln.mpg.de/GABI-Kat/>.

FEATURES Location/Qualifiers

1..32

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Columbia 0"

/db_xref="taxon:3702"

/clone="GK-257B04-014921"

/clone_lib="Arabidopsis thaliana T-DNA insertion lines"

/note="PCR was performed on DNA from Arabidopsis thaliana plants (T1) which were transformed with the T-DNA from vector pAC161. The lines contain one or more T-DNA from insertions. The DNA fragment(s) resulting from the PCR were directly sequenced to determine the genomic sequence flanking the insertion. Sequences displaying significant similarity to the A. thaliana nuclear genome sequences were processed for submission. T-DNA derived sequences were removed"

0 a 6 c 0 g 26 t

BASE COUNT 0 a 6 c 0 g 26 t

ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 32;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTTTT 10

|||||

Db 10 CTCTCTTTT 19

RESULT 7

BZ763249

LOCUS BZ763249 33 bp DNA linear GSS 13-MAR-2003

DEFINITION SALK_115724.25.30.n Arabidopsis thaliana T-DNA insertion lines Arabidopsis thaliana genomic clone SALK_115724.25.30.n, genomic survey sequence.

ACCESSION BZ763249

VERSION BZ763249.1 GI:28935802

KEYWORDS GSS

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM Arabidopsis thaliana

REFERENCE 1 (bases 1 to 33)

AUTHORS Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R., Gadrinab, C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prehn, L., Shinn, P., Zimmerman, J. and Ecker, J.R.

TITLE A Sequence-Indexed Library of Insertion Mutations in the Arabidopsis Genome

JOURNAL Unpublished

COMMENT Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGNAL)

TITLE A pipeline for automated high-throughput generation of ESTs (flanking sequence tags) from Arabidopsis thaliana T-DNA transformed lines

JOURNAL Unpublished

REFERENCE 2

AUTHORS Rosso, M., Strizhov, N., Li, Y., Reiss, B., Dekker, K. and Weissshaar, B.

TITLE A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat) for flanking sequence tag based reverse genetics

JOURNAL Unpublished

REFERENCE 3 (bases 1 to 32)

AUTHORS Rosso, M., Li, Y., Strizhov, N. and Weissshaar, B.

TITLE Direct Submission

JOURNAL Submitted (21-OCT-2002) Weissshaar B., Max-Planck-Institut fuer Zuechtungsforschung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany

COMMENT This sequence is recovered from the left border of the T-DNA. It indicates an insertion close to or within gene At5g42900. The sequences are generated at the MPI for Plant Breeding Research in the context of the GABI-Kat project. GABI-Kat is part of the German Plant Genomics program designated 'GABI'. Information on line availability can be found at: <http://www.mpiz-koeln.mpg.de/GABI-Kat/>.

FEATURES Location/Qualifiers

1..33

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Columbia 0"

/db_xref="taxon:3702"

/clone="SALK_115724.25.30.n"

/clone_lib="Arabidopsis thaliana T-DNA insertion lines"

/note="PCR was performed on Arabidopsis thaliana lines each of which contains one or more T-DNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at http://signal.salk.edu/tdna_protocols.html"

9 a 7 c 3 g 14 t

BASE COUNT 9 a 7 c 3 g 14 t

ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 33;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTTTT 10

|||||

Db 10 CTCTCTTTT 19

RESULT 8

AU256102/c

LOCUS AU256102 34 bp mRNA linear EST 25-APR-2002

DEFINITION AU256102 3'-directed mouse cDNA library Mus musculus cDNA clone BED0007468 3', mRNA sequence.

ACCESSION AU256102

VERSION AU256102.1 GI:20319468

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 34)

AUTHORS Kato, K. and Matoba, R.

TITLE Generation of expressed sequence tags from mouse brain

JOURNAL Unpublished

COMMENT Contact: Kikuya Kato
Graduate School of Biological Sciences
Nara Institute of Science and Technology
8916-5 Takayama, Ikoma, Nara 630-0101, Japan
Tel: 81-743-72-5581
Fax: 81-743-72-5589
Email: kkatobs.aistc-nara.ac.jp, /BED/index.html.
URL: <http://love2.aist-nara.ac.jp/BED/index.html>.

FEATURES Location/Qualifiers

1..34

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="BED0007468"

/tissue type="brain"

/clone_lib="3'-directed mouse cDNA library"

14 a 5 c 7 g 8 t

BASE COUNT 14 a 5 c 7 g 8 t

ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 34;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of TDNA.

Class: TDNA tagged.

FEATURES Location/Qualifiers

source

1..33

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Columbia 0"

/db_xref="taxon:3702"

/clone="SALK_115724.25.30.n"

/clone_lib="Arabidopsis thaliana T-DNA insertion lines"

/note="PCR was performed on Arabidopsis thaliana lines each of which contains one or more T-DNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at http://signal.salk.edu/tdna_protocols.html"

BASE COUNT 9 a 7 c 3 g 14 t

ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 33;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTTTT 10

|||||

Db 10 CTCTCTTTT 19

RESULT 8

AU256102/c

LOCUS

AU256102 3'-directed mouse cDNA library Mus musculus cDNA clone

DEFINITION BED0007468 3', mRNA sequence.

ACCESSION AU256102

VERSION AU256102.1 GI:20319468

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 34)

AUTHORS Kato, K. and Matoba, R.

TITLE Generation of expressed sequence tags from mouse brain

JOURNAL Unpublished

COMMENT Contact: Kikuya Kato

Graduate School of Biological Sciences

Nara Institute of Science and Technology

8916-5 Takayama, Ikoma, Nara 630-0101, Japan

Tel: 81-743-72-5581

Fax: 81-743-72-5589

Email: kkatobs.aistc-nara.ac.jp, /BED/index.html.

URL: <http://love2.aist-nara.ac.jp/BED/index.html>.

FEATURES Location/Qualifiers

1..34

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="BED0007468"

/tissue type="brain"

/clone_lib="3'-directed mouse cDNA library"

14 a 5 c 7 g 8 t

BASE COUNT 14 a 5 c 7 g 8 t

ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 34;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
 Db 20 CTTCTCTTTT 11

RESULT 9
 AZ429862 37 bp DNA linear GSS 03-OCT-2000
 LOCUS
 DEFINITION IM0214105F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0214105 F, genomic survey sequence.
 ACCESSION AZ429862
 VERSION AZ429862.1 GI:10553875
 KEYWORDS GSS.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 37)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
 ,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
 and Wright,D.,Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 JOURNAL Unpublished
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0214 row: I column: 05
 Seq primer: CGTTGTAACGACGCCAGT
 Class: plasmid ends
 High quality sequence stop: 37.
 Location/Qualifiers
 1..37
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0214105"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /note="Vector: PWD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adapted DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pWD42 (gi|4732114|gb|AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adapted mouse DNA was annealed to
 adapted vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

FEATURES
 source

1..37
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0214105"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /note="Vector: PWD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adapted DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pWD42 (gi|4732114|gb|AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adapted mouse DNA was annealed to
 adapted vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

BASE COUNT
 ORIGIN

8 a 9 c 6 g 14 t

Query Match 100.0%; Score 10; DB 28; Length 37;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
 Db 21 CTTCTCTTTT 30

RESULT 10
 AZ514585 37 bp DNA linear GSS 05-OCT-2000
 LOCUS
 DEFINITION IM0361B16F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0361B16 F, genomic survey sequence.
 ACCESSION AZ514585
 VERSION AZ514585.1 GI:10695817
 KEYWORDS GSS
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 37)
 AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
 ,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
 and Wright,D.,Weiss,R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 JOURNAL Unpublished
 COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0361 row: B column: 16
 Seq primer: CGTTGTAACGACGCCAGT
 Class: plasmid ends
 High quality sequence stop: 37.
 Location/Qualifiers
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 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0361B16"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /note="Vector: PWD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adapted DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pWD42 (gi|4732114|gb|AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adapted mouse DNA was annealed to
 adapted vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

FEATURES
 source

1..37
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0361B16"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /note="Vector: PWD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adapted DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pWD42 (gi|4732114|gb|AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adapted mouse DNA was annealed to
 adapted vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

BASE COUNT 3 a 8 c 2 g 24 t

Query Match 100.0%; Score 10; DB 28; Length 37;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||
 Db 12 CTTCTCTTTT 21

RESULT 11
 AZ613373
 LOCUS
 DEFINITION
 1M0441M12R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0441M12 R, genomic survey sequence.

ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
 Mus musculus (house mouse)

REFERENCE
 AUTHORS
 Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
 M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
 and Wright,D., Weiss,R.

TITLE
 Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts

JOURNAL
 COMMENT
 Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
 84112, USA

FEATURES
 source
 1..39
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0441M12"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /note="Vector: PWD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adaptor DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pWD42 (gi|4732114|gb|AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

BASE COUNT
 ORIGIN
 8 a 8 c 2 g 21 t

Query Match 100.0%; Score 10; DB 28; Length 39;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||
 Db 2 CTTCTCTTTT 11

RESULT 12
 AA916182/c
 LOCUS
 DEFINITION
 AA916182 40 bp mRNA linear EST 14-APR-1998
 Q334B06.s1 NCI CGAP Br7 Homo sapiens cDNA clone IMAGE:1441715 3',
 similar to TR:Q33575 Q33575 NADH DEHYDROGENASE SUBUNIT 4. ;, mRNA
 sequence.

ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
 Homo sapiens (human)

REFERENCE
 AUTHORS
 NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index

JOURNAL
 COMMENT
 Contact: Robert Strausberg, Ph.D.
 Email: cgapsb-remail.nih.gov
 unknown library type
 Trace considered overall poor quality
 Seq primer: -40m13 fwd. ET from Amersham
 High quality sequence stop: 1.
 Location/Qualifiers

FEATURES
 source
 1..40
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:1441715"
 /lab_host="DH10B"
 /clone_lib="NCI-CGAP Br7"
 /note="Organ: breast; Vector: pCMV-SPORT4; Site 1: SalI;
 Site 2: NotI; Cloned unidirectionally. Primer: Oligo dT.
 Average insert size 1.2 kb. Life Technologies catalog
 #:10985-018"

BASE COUNT 26 a 12 g 2 t

Query Match 100.0%; Score 10; DB 9; Length 40;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||
 Db 18 CTTCTCTTTT 9

RESULT 13
 AZ375959/c
 LOCUS
 DEFINITION
 AZ375959 41 bp DNA linear GSS 02-OCT-2000
 1M0129A12R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0129A12 R, genomic survey sequence.

ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
 Mus musculus (house mouse)

REFERENCE
 AUTHORS
 Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
 Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
 M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
 and Wright,D., Weiss,R.

TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts
JOURNAL Unpublished
COMMENT Contact: Robert B. Weiss
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: dunnegenetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0129 row: A column: 12
 Seq primer: CACACAGGAACAGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 41.

FEATURES source
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 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0129A12"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /note="Vector: PWB42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt ended-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (G1.4732114|GB|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 22 a 4 c 7 g 8 t
ORIGIN
 Query Match 100.0%; Score 10; DB 28; Length 41;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
Db 25 CTTCTCTTTT 16

RESULT 14
EX120830/c
LOCUS BX120830 42 bp DNA linear GSS 13-MAR-2003
DEFINITION Danio rerio genomic clone DKEY-61D1, genomic survey sequence.
ACCESSION BX120830
VERSION BX120830.1 GI:27951749
KEYWORDS GSS.
SOURCE Danio rerio (zebrafish)
ORGANISM Danio rerio
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes; Cyprinidae; Danio.
 1 (bases 1 to 42)
 Humphray,S.J., Huckle,E. and Durham,J.L.
 Direct Submission
 Submitted (13-MAR-2003) The Sanger Institute, Wellcome Trust Genome Campus, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Unpublished
 This sequence was generated from the SP6 end of BAC 64G19. 64G19 is part of the Daniokey BAC library created by R. Plasterk and N.V. Keygene. Further details:
 http://www.sanger.ac.uk/projects/D_rerio/.

FEATURES source
 1..42
 /organism="Danio rerio"
 /mol_type="genomic DNA"
 /db_xref="taxon:7955"
 /clone="DKEY-61D1"
 /tissue type="Testis"
 /note="vector pIndigoBAC-536"
 19 a 9 c 9 g 0 t

BASE COUNT 19 a 9 c 9 g 0 t
ORIGIN
 Query Match 100.0%; Score 10; DB 29; Length 42;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
Db 10 CTTCTCTTTT 1

RESULT 16
CC050354/c
LOCUS CC050354 45 bp DNA linear GSS 01-APR-2003
DEFINITION OIS-536-6-lto6-G01 UniformMu Mutail Library Zea mays genomic clone OIS-536-6-lto6-G01, genomic survey sequence.

COMMENT Campus, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Unpublished
 This sequence was generated from the T7 end of BAC 61D1. 61D1 is part of the Daniokey BAC Library created by R. Plasterk and N.V. Keygene. Further details:
 http://www.sanger.ac.uk/projects/D_rerio/.

FEATURES source
 1..42
 /organism="Danio rerio"
 /mol_type="genomic DNA"
 /db_xref="taxon:7955"
 /clone="DKEY-61D1"
 /tissue type="Testis"
 /note="vector pIndigoBAC-536"
 29 a 0 c 13 g 0 t

BASE COUNT 29 a 0 c 13 g 0 t
ORIGIN
 Query Match 100.0%; Score 10; DB 29; Length 42;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
Db 27 CTTCTCTTTT 18

RESULT 15
EX124544/c
LOCUS BX124544 42 bp DNA linear GSS 28-JAN-2003
DEFINITION Danio rerio genomic clone DKEY-64G19, genomic survey sequence.
ACCESSION BX124544
VERSION BX124544.1 GI:27955482
KEYWORDS GSS.
SOURCE Danio rerio (zebrafish)
ORGANISM Danio rerio
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes; Cyprinidae; Danio.
 1 (bases 1 to 42)
 Humphray,S.J., Huckle,E. and Durham,J.L.
 Direct Submission
 Submitted (27-JAN-2003) The Sanger Institute, Wellcome Trust Genome Campus, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Unpublished
 This sequence was generated from the SP6 end of BAC 64G19. 64G19 is part of the Daniokey BAC library created by R. Plasterk and N.V. Keygene. Further details:
 http://www.sanger.ac.uk/projects/D_rerio/.

FEATURES source
 1..42
 /organism="Danio rerio"
 /mol_type="genomic DNA"
 /db_xref="taxon:7955"
 /clone="DKEY-64G19"
 /tissue type="Testis"
 /note="vector pIndigoBAC-536"
 19 a 9 c 9 g 0 t

BASE COUNT 19 a 9 c 9 g 0 t
ORIGIN
 Query Match 100.0%; Score 10; DB 29; Length 42;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
Db 10 CTTCTCTTTT 1

RESULT 16
CC050354/c
LOCUS CC050354 45 bp DNA linear GSS 01-APR-2003
DEFINITION OIS-536-6-lto6-G01 UniformMu Mutail Library Zea mays genomic clone OIS-536-6-lto6-G01, genomic survey sequence.

<p>CC050354 VERSION CC050354.1 GI:29465245 KEYWORDS GSS. SOURCE Zea mays ORGANISM Zea mays</p>	<p>Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD clade; Panicoideae; Andropogoneae; Zea.</p>	<p>Lathaw,S., Tan,B.-C., Settles,A.M. and McCarty,D.R. Sequence tagged transposon insertions from the UniformMu maize population Unpublished Contact: Donald R. McCarty Plant Molecular and Cellular Biology Program University of Florida PO 110690 Gainesville, FL 32611-0690, USA Tel: 352-392-1928 x322 Email: drmc@ufl.edu Sequence flanking probable Mu insertion site in UniformMu line: 01S-536-6 Class: transposon insertion site.</p>	<p>1. .45 Location/Qualifiers /organism="Zea mays" /mol_type="genomic DNA" /strain="W22 (ACR, bz1-m9)" /cultivar="UniformMu" /db_xref="taxon:4577" /clone="01S-536-6-lto6-G01" /clone_lib="UniformMu Mutail Library" /notes="Vector: TOPO-PCR4; DNA flanking Mu transposon insertions in Mu inactive lines were extracted from the UniformMu maize population by the thermo asymmetric interlaced PCR (TAIL) protocol using primers specific for the Mu terminal inverted repeat and a set of 16 arbitrary primers. Amplicons were size enriched using Sepharose 400 spin columns and cloned into the TOPO PCR4 vector."</p>	<p>BASE COUNT 17 a 6 c 16 g 6 t ORIGIN</p>	<p>Query Match 100.0%; Score 10; DB 29; Length 45; Best Local Similarity 100.0%; Pred. No. 1.6e+05; Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;</p>	<p>Qy 1 CTTCTCTTTT 10 Db 20 CTTCTCTTTT 11</p>	<p>RESULT 17 BX120970/c LOCUS BX120970 45 bp DNA linear GSS 28-JAN-2003 DEFINITION Danio rerio genomic clone DKEI-68A12, genomic survey sequence. ACCESSION BX120970 VERSION BX120970.1 GI:27951891 KEYWORDS GSS. SOURCE Danio rerio (zebrafish) ORGANISM Danio rerio Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes; Cyprinidae; Danio. 1 (bases 1 to 45) Humphray,S.J., Huckle,E. and Durham,J.L. Direct Submission Submitted (27-JAN-2003) The Sanger Institute, Wellcome Trust Genome Campus, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Unpublished This sequence was generated from the SP6 end of BAC 68A12. 68A12 is part of the Dantokoy BAC Library created by R. Plasterk and N.V. Keygene. Further details: http://www.sanger.ac.uk/Projects/D_rerio/ Location/Qualifiers</p>	<p>Accession CC050354 Version CC050354.1 GI:29465245 Organism Zea mays Source Zea mays Reference Lathaw,S., Tan,B.-C., Settles,A.M. and McCarty,D.R. Authors Lathaw,S., Tan,B.-C., Settles,A.M. and McCarty,D.R. Title Sequence tagged transposon insertions from the UniformMu maize population Journal Unpublished Comment Contact: Donald R. McCarty Plant Molecular and Cellular Biology Program University of Florida PO 110690 Gainesville, FL 32611-0690, USA Tel: 352-392-1928 x322 Email: drmc@ufl.edu Sequence flanking probable Mu insertion site in UniformMu line: 01S-536-6 Class: transposon insertion site.</p>	<p>1. .45 Location/Qualifiers /organism="Danio rerio" /mol_type="genomic DNA" /db_xref="taxon:7955" /clone="DKEI-68A12" /tissue type="Testis" /note="vector pIndigoBAC-536" 28 a 2 c 15 g 0 t</p>	<p>BASE COUNT 28 a 2 c 15 g 0 t ORIGIN</p>	<p>Query Match 100.0%; Score 10; DB 29; Length 45; Best Local Similarity 100.0%; Pred. No. 1.6e+05; Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;</p>	<p>Qy 1 CTTCTCTTTT 10 Db 21 CTTCTCTTTT 12</p>	<p>RESULT 18 AA922976 LOCUS AA922976 49 bp mRNA linear EST 21-APR-1998 DEFINITION ok77e06.sl NCI CGAP GC4 Homo sapiens cDNA clone IMAGE:1520002 3' similar to TR:Q39614 Q39614 PROLINE-RICH PROTEIN.; mRNA sequence. ACCESSION AA922976 VERSION AA922976.1 GI:3070285 KEYWORDS EST. SOURCE Homo sapiens (human) ORGANISM Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 49) NCI-CGAP http://www.ncbi.nlm.nih.gov/hcicgap. National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index Unpublished Contact: Robert Strausberg, Ph.D. Email: cgabpsr@mail.nih.gov Tissue Procurement: Christopher A. Moskaluk, M.D., Ph.D., Michael Emmert-Buck, M.D., Ph.D. CDNA Library Preparation: M. Bento Soares, Ph.D. CDNA Library Arrayed by: Greg Lennon, Ph.D. DNA Sequencing by: Washington University Genome Sequencing Center Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www.bio.llnl.gov/bbrp/image/image.html</p>	<p>Trace considered overall poor quality Seq primer: -40ml3 fwd. ET from Amersham High quality sequence stop: 1. Location/Qualifiers 1. .49 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="taxon:9606" /clone="IMAGE:1520002" /tissue_type="pooled germ cell tumors" /lab host="DH10B" /clone_lib="NCI CGAP GC4" /notes="Vector: p7T3D-Pac (Pharmacia) with a modified polylinker; 1st strand cDNA was prepared from 3 pooled germ cell tumors, and was then primed with a Not I - oligo(dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified p7T3 vector. Library is normalized. Library was constructed by Bento Soares and M. Fatima Bonaldo." 10 a 23 c 0 g 16 t</p>	<p>BASE COUNT 10 a 23 c 0 g 16 t ORIGIN</p>	<p>Query Match 100.0%; Score 10; DB 9; Length 49; Best Local Similarity 100.0%; Pred. No. 1.6e+05; Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;</p>	<p>Accession CC050354 Version CC050354.1 GI:29465245 Organism Zea mays Source Zea mays Reference Lathaw,S., Tan,B.-C., Settles,A.M. and McCarty,D.R. Authors Lathaw,S., Tan,B.-C., Settles,A.M. and McCarty,D.R. Title Sequence tagged transposon insertions from the UniformMu maize population Journal Unpublished Comment Contact: Donald R. McCarty Plant Molecular and Cellular Biology Program University of Florida PO 110690 Gainesville, FL 32611-0690, USA Tel: 352-392-1928 x322 Email: drmc@ufl.edu Sequence flanking probable Mu insertion site in UniformMu line: 01S-536-6 Class: transposon insertion site.</p>	<p>1. .45 Location/Qualifiers /organism="Zea mays" /mol_type="genomic DNA" /strain="W22 (ACR, bz1-m9)" /cultivar="UniformMu" /db_xref="taxon:4577" /clone="01S-536-6-lto6-G01" /clone_lib="UniformMu Mutail Library" /notes="Vector: TOPO-PCR4; DNA flanking Mu transposon insertions in Mu inactive lines were extracted from the UniformMu maize population by the thermo asymmetric interlaced PCR (TAIL) protocol using primers specific for the Mu terminal inverted repeat and a set of 16 arbitrary primers. Amplicons were size enriched using Sepharose 400 spin columns and cloned into the TOPO PCR4 vector."</p>	<p>BASE COUNT 17 a 6 c 16 g 6 t ORIGIN</p>	<p>Query Match 100.0%; Score 10; DB 29; Length 45; Best Local Similarity 100.0%; Pred. No. 1.6e+05; Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;</p>	<p>Qy 1 CTTCTCTTTT 10 Db 20 CTTCTCTTTT 11</p>	<p>RESULT 17 BX120970/c LOCUS BX120970 45 bp DNA linear GSS 28-JAN-2003 DEFINITION Danio rerio genomic clone DKEI-68A12, genomic survey sequence. ACCESSION BX120970 VERSION BX120970.1 GI:27951891 KEYWORDS GSS. SOURCE Danio rerio (zebrafish) ORGANISM Danio rerio Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes; Cyprinidae; Danio. 1 (bases 1 to 45) Humphray,S.J., Huckle,E. and Durham,J.L. Direct Submission Submitted (27-JAN-2003) The Sanger Institute, Wellcome Trust Genome Campus, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk Unpublished This sequence was generated from the SP6 end of BAC 68A12. 68A12 is part of the Dantokoy BAC Library created by R. Plasterk and N.V. Keygene. Further details: http://www.sanger.ac.uk/Projects/D_rerio/ Location/Qualifiers</p>	<p>Accession CC050354 Version CC050354.1 GI:29465245 Organism Zea mays Source Zea mays Reference Lathaw,S., Tan,B.-C., Settles,A.M. and McCarty,D.R. Authors Lathaw,S., Tan,B.-C., Settles,A.M. and McCarty,D.R. Title Sequence tagged transposon insertions from the UniformMu maize population Journal Unpublished Comment Contact: Donald R. McCarty Plant Molecular and Cellular Biology Program University of Florida PO 110690 Gainesville, FL 32611-0690, USA Tel: 352-392-1928 x322 Email: drmc@ufl.edu Sequence flanking probable Mu insertion site in UniformMu line: 01S-536-6 Class: transposon insertion site.</p>	<p>1. .45 Location/Qualifiers /organism="Zea mays" /mol_type="genomic DNA" /strain="W22 (ACR, bz1-m9)" /cultivar="UniformMu" /db_xref="taxon:4577" /clone="01S-536-6-lto6-G01" /clone_lib="UniformMu Mutail Library" /notes="Vector: TOPO-PCR4; DNA flanking Mu transposon insertions in Mu inactive lines were extracted from the UniformMu maize population by the thermo asymmetric interlaced PCR (TAIL) protocol using primers specific for the Mu terminal inverted repeat and a set of 16 arbitrary primers. Amplicons were size enriched using Sepharose 400 spin columns and cloned into the TOPO PCR4 vector."</p>	<p>BASE COUNT 17 a 6 c 16 g 6 t ORIGIN</p>	<p>Query Match 100.0%; Score 10; DB 29; Length 45; Best Local Similarity 100.0%; Pred. No. 1.6e+05; Matches </p>
--	---	---	--	--	---	---	---	---	---	--	---	---	--	--	---	--	---	--	--	---	---	---	---	--	--	--

```

QY      1 CTTCTCTTTT 10
Db      1 CTTCTCTTTT 10

RESULT 19
A1564984
LOCUS   49 bp  mRNA  linear  EST 13-MAY-1999
DEFINITION
tq53b02.x1 NCI CGAP Utl1 Homo sapiens cDNA clone IMAGE:2212491 3'
similar to WP.F59E12.9 CELL1534 ;contains element MER22 repetitive
element ;, mRNA sequence.

ACCESSION
A1564984
VERSION 1
KEYWORDS
SOURCE  Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 49)
REFERENCE
A1564984
AUTHORS
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL
Unpublished
COMMENT
Contact: Robert Strausberg, Ph.D.
Email: cgabps-remail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

Trace considered overall poor quality
Insert length: 400 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 1
POLYA=No.

FEATURES             Location/Qualifiers
     source           1..49
     /organism="Homo sapiens"
     /mol_type="mRNA"
     /db_xref="taxon:9606"
     /clone="IMAGE:2212491"
     /tissue type="well-differentiated endometrial
adenocarcinoma, 7 pooled tumors"
     /lab_host="DH10B"
     /clone lib="NCI CGAP Utl1"
     /notes="Organ: uterus; Vector: pCMV-SPORT6; Site 1: SalI;
Site 2: NotI; Cloned unidirectionally. Primer: Oligo dt.
Average insert size 1.75 kb. Life Technologies catalog #:
11538-014"

BASE COUNT      0 a 26 c 0 g 23 t
ORIGIN
Query Match      100.0%; Score 10; DB 9; Length 49;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      9 CTTCTCTTTT 18

RESULT 20
BZ357069
LOCUS   49 bp  DNA  linear  GSS 14-NOV-2002
DEFINITION
SALK_130198.43.95.x Arabidopsis thaliana TDNA insertion lines
Arabidopsis thaliana genomic clone SALK_130198.43.95.x, genomic
survey sequence.

ACCESSION
BZ357069

```

```

VERSION BZ357069.1 GI:24948818
KEYWORDS GSS.
SOURCE Arabidopsis thaliana (chale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosid II; Brassicales; Brassicaceae; Arabidopsi.
1 (bases 1 to 49)
REFERENCE
A102691
AUTHORS
Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab
,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.,
Zimmerman,J. and Ecker,J.R.
TITLE
A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
JOURNAL
Unpublished
COMMENT
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGnAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: eckersalk.edu
This is single pass sequence recovered from the left border of
TDNA.
Class: TDNA tagged.
Location/Qualifiers
     1..49
     /organism="Arabidopsis thaliana"
     /mol_type="genomic DNA"
     /strain="Columbia 0"
     /db_xref="taxon:3702"
     /clone="SALK_130198.43.95.x"
     /clone lib="Arabidopsis thaliana TDNA insertion lines"
     /note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"

BASE COUNT      12 a 13 c 5 g 19 t
ORIGIN
Query Match      100.0%; Score 10; DB 29; Length 49;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      12 CTTCTCTTTT 21

RESULT 21
A102691/c
LOCUS   50 bp  mRNA  linear  EST 30-AUG-2001
DEFINITION
A102691 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
HMA230056, mRNA sequence.

ACCESSION
A102691
VERSION A102691.1 GI:13552212
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 50)
REFERENCE
A102691
AUTHORS
Suzuki,Y., Taira,H., Tsunoda,T., Mizushima-Sugano,J., Sese,J., Hata
,H., Ota,T., Isogai,T., Tanaka,T., Morishita,S., Okubo,K., Sakaki
,Y., Nakamura,Y., Suyama,A. and Sugano,S.
TITLE
Diverse transcriptional initiation revealed by fine, large-scale
mapping of mRNA start sites
JOURNAL
EMBO Rep. 2 (5), 388-393 (2001)
MEDLINE 21270072
PUBMED 11375929
COMMENT
Contact: Yutaka Suzuki
Department of Virology

```

Institute of Medical Science, University of Tokyo
4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
Email: yasuku@ims.u-tokyo.ac.jp
Suzuki, Y., Yoshitomo-Nakagawa, K., Maruyama, K., Suyama, A. and Sugano
S. Construction and characterization of a full length-enriched and
a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

FEATURES

source

1. .50
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="HEM230056"
/clone_lib="Sugano Homo sapiens cDNA library"

BASE COUNT

13 a 10 c 17 g 10 t

Query Match

Best Local Similarity 100.0%; Score 10; DB 9; Length 50;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY

1 CTTCTCTTTT 10

|||||

Db

28 CTTCTCTTTT 19

RESULT 22

BZ770334

LOCUS

DEFINITION SALK_143268.36.90.x Arabidopsis thaliana TDNA insertion lines
Arabidopsis thaliana genomic clone SALK_143268.36.90.x, genomic
survey sequence.

ACCESSION

BZ770334

VERSION

BZ770334.1 GI:28944018

KEYWORDS

SOURCE

ORGANISM

Arabidopsis thaliana (thale cress)

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosid II; Brassicales; Brassicaceae; Arabidopsids.

REFERENCE

AUTHORS

Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R., Gadriab
C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prednis, L., Shinn, P.,
Zimmerman, J., and Ecker, J.R.

A Sequence-Indexed Library of Insertion Mutations in the

Arabidopsis Genome

Unpublished

Contact: Joseph R. Ecker

Salk Institute Genomic Analysis Laboratory (SIGAL)

The Salk Institute for Biological Studies

10010 N. Torrey Pines Road, La Jolla, CA 92037, USA

Tel: 858 453 4100 x1752

Fax: 858 558 6379

Email: ecker@salk.edu

This is single pass sequence recovered from the left border of

TDNA.

Class: TDNA tagged.

Location/Qualifiers

1. .51

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Columbia 0"

/db_xref="taxon:3702"

/clone="SALK_143268.36.90.x"

/clone_lib="Arabidopsis thaliana TDNA insertion lines"

/note="PCR was performed on Arabidopsis thaliana lines

each of which contains one or more TDNA insertion

elements. The resultant fragment for each line was

directly sequenced to determine the genomic sequence at

the site of insertion. Details of the protocols used can

be found at http://signal.salk.edu/tdna_protocols.html

14 a 10 c 5 g 22 t

BASE COUNT

ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 51;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY

1 CTTCTCTTTT 10

|||||

Db

17 CTTCTCTTTT 26

RESULT 23

AB082596

LOCUS

DEFINITION

Drosophila melanogaster DNA, clone:1(2)SH2 1592, genomic survey

sequence.

ACCESSION

AB082596

VERSION

AB082596.1 GI:24416782

KEYWORDS

SOURCE

ORGANISM

Drosophila melanogaster (fruit fly)

Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;

Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;

Ephydroidea; Drosophilidae; Drosophila.

REFERENCE

AUTHORS

Oh, S., Kingsley, T., Shin, H., Zheng, Z., Chen, H. and Hou, S.

Functional Genomics: A P element-mediated gene disruption in

Drosophila

Unpublished

2 (bases 1 to 51)

Oh, S., Kingsley, T., Shin, H., Zheng, Z., Chen, H. and Hou, S.

Direct Submission

Submitted (26-MAR-2002) Suwan Oh, NCI-FCRDC, Lab. Of Immunobiology;

1050 Boyles st., Frederick, Maryland 21702, USA

(E-mail: ohsuwan@mail.nih.gov, Tel: 1-301-846-7314,

Fax: 1-301-846-6145)

Location/Qualifiers

1. .51

/organism="Drosophila melanogaster"

/mol_type="genomic DNA"

/db_xref="taxon:7227"

/clone="1(2)SH2 1592"

6 a 19 c 8 g 18 t

BASE COUNT

ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 51;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY

1 CTTCTCTTTT 10

|||||

Db

28 CTTCTCTTTT 37

|||||

RESULT 24

BQ667496/c

LOCUS

DEFINITION

Ancyllostoma caninum cDNA 5', mRNA sequence.

ACCESSION

BQ667496

VERSION

BQ667496.1 GI:21809178

KEYWORDS

SOURCE

ORGANISM

Ancyllostoma caninum (dog hookworm)

Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Strongylida;

Ancyllostomatoidea; Ancyllostomatidae; Ancyllostomatinae; Ancyllostoma.

1 (bases 1 to 52)

McCarte, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J., Wylie, T.

, Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y.

, Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Tsagarishvili, R.

, Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Steptoe

, M., Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S.

, Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and

Wilson, R.

The Washington Univ. Nematode EST Project, 1999

EST.

Ancyllostoma caninum

Ancyllostoma caninum

52 bp mRNA linear EST 15-JUL-2002

pb62c12.y1 Anc caninum i3 serum stim pAMPI v1 Chiapelli McCarter

Ancyllostoma caninum cDNA 5', mRNA sequence.

QY

1 CTTCTCTTTT 10

|||||

Db

28 CTTCTCTTTT 37

|||||

JOURNAL
COMMENT

Unpublished
Contact: McCarter JP
The Washington Univ. Nematode EST Project, 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wustl.wustl.edu
The library was constructed by Brandi Chiapelli and Dr. James McCarter (bchiapell@wustl.edu & jmcarter@wustl.edu) at Washington University, St. Louis. DNA Sequencing by: Washington University Genome Sequencing Center St. Louis. Nematodes were provided by Dr. Prema Arasu of North Carolina State University. Putative full length read
The vector to vector length is 53.

FEATURES

source

1. 52
/organism="Ancylostoma caninum"
/mol_type="mRNA"
/db_xref="taxon:29170"
/dev_stage="serum stimulated L3"
/lab_host="DH10B"
/clone_lib="Anc caninum L3 serum stim pAMP1 v1 Chiapelli McCarter"
/note="Vector: pAMP1 (Gibco); Site_1: NotI; Site_2: SalI; the library was constructed by Brandi Chiapelli and Dr. James McCarter at Washington University, St. Louis. The cDNA was made by using Dynabead oligo-dT priming (Dyna) PCR based library using a modified protocol from the SMART PCR cDNA Synthesis Kit from Clontech. Directionally cloned into the UDG sites of pAMP1. Nematodes were provided by Dr. Prema Arasu of North Carolina State University."

BASE COUNT 32 a 3 c 13 g 4 t
ORIGIN

Query Match 100.0%; Score 10; DB 13; Length 52;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 44 CTTCTCTTTT 35

RESULT 25
AZ327146/c

LOCUS 52 bp DNA linear GSS 29-SEP-2000
DEFINITION 1M0050016F Mouse 10kb plasmid UUGC1M library Mus musculus genomic clone UUGC1M0050016 F, genomic survey sequence.

ACCESSION AZ327146.1 GI:10385604

VERSION GSS.

KEYWORDS Mus musculus (house mouse)

ORGANISM

Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 52)

Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D.,Weiss,R.

Mouse whole genome scaffolding with paired end reads from 10kb

plasmid inserts

TITLE

JOURNAL

COMMENT

Unpublished
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA

Tel: 801 585 5606

Fax: 801 585 7177

Email: dduunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0050 row: 0 column: 16

Seq primer: CGTTGTAAACGACGCGCCAGT

Class: plasmid ends

High quality sequence stop: 52.

Location/Qualifiers

source

1. 52
/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC1M0050016"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, Tl-resistant, F-"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/note="Vector: pMD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pMD42 (GI:4732114|gb|AF129072.1), a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT 26 a 7 c 9 g 10 t
ORIGIN

Query Match 100.0%; Score 10; DB 28; Length 52;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 32 CTTCTCTTTT 23

RESULT 26

BE569086/c

LOCUS

DEFINITION 601339390F2 NIH_MGC_53 Homo sapiens cDNA clone IMAGE:3681382 5', mRNA sequence.

ACCESSION BE569086

VERSION BE569086.1 GI:9812806

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 55)

NIH-MGC http://mgs.nci.nih.gov/.

Unpublished

National Institutes of Health, Mammalian Gene Collection (MGC)

Contact: Robert Strausberg, Ph.D.

Email: cgabbs@mail.nih.gov

Tissue Procurement: ATCC

cDNA Library Preparation: CLONTECH Laboratories, Inc.

DNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov

Plate: LLC363 row: d column: 23.

Location/Qualifiers

1. 52

FEATURES

source


```

/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:3681382"
/tissue_type="carcinoma, cell line"
/lab_host="DH10B (T1 phage-resistant)"
/clone_lib="NIH MGC 53"
/note="Organ: bladder; Vector: pDNR-LIB (Clontech);
Site 1: SfiI (ggcgctcgcc); Site 2: SfiI (ggccattatggcc
); Double-stranded cDNA was prepared from cell line RNA.
5' and 3' adaptors were used in cloning as follows: 5'
adaptor sequence: 5'-CAGGCCATTATGCC-3' and 3' adaptor
sequence: 5'-ATTCTAGAGCGGCGCGGCACATG-dt(30)BN-3'
(where B = A, C, or G and N = A, C, G, or T). Average
insert size 1.55 kb (range 0.9-4.0 kb). 15/15 colonies
contained inserts by PCR. This library was enriched for
full-length clones and was constructed by Clontech
Laboratories (Palo Alto, CA)."
BASE COUNT      40 a      1 c      11 g
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      48 CTTCTCTTTT 39

RESULT 27
BH908756      55 bp      DNA      linear      GSS 04-SEP-2002
LOCUS
DEFINITION
SALK_050419.22.05.x Arabidopsis thaliana TDNA insertion lines
Arabidopsis thaliana genomic clone SALK_050419.22.05.x, genomic
survey sequence.
ACCESSION      BH908756
VERSION
KEYWORDS
SOURCE
ORGANISM
Arabidopsis thaliana (thale cress)
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 55)
REFERENCE
AUTHORS
Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab
,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.,
Zimmerman,J. and Ecker,J.R.
A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
Unpublished
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGnAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of
TDNA.
FEATURES
Class: TDNA tagged.
Location/Qualifiers
1..55
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="SALK_050419.22.05.x"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"
BASE COUNT      19 a      5 c      12 g      20 t
ORIGIN
Query Match      100.0%; Score 10; DB 28; Length 56;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      53 CTTCTCTTTT 44

RESULT 29
BH222307
LOCUS

```

```

the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"
BASE COUNT      9 a      15 c      3 g      28 t
ORIGIN
Query Match      100.0%; Score 10; DB 28; Length 55;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      42 CTTCTCTTTT 51

RESULT 28
BH864698/c      56 bp      DNA      linear      GSS 05-AUG-2002
LOCUS
DEFINITION
SALK_096733 Arabidopsis thaliana TDNA insertion lines Arabidopsis
thaliana genomic clone SALK_096733, genomic survey sequence.
ACCESSION      BH864698
VERSION
KEYWORDS
SOURCE
ORGANISM
Arabidopsis thaliana (thale cress)
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 56)
REFERENCE
AUTHORS
Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab
,C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.,
Zimmerman,J. and Ecker,J.R.
A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
Unpublished
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGnAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of
TDNA.
FEATURES
Class: TDNA tagged.
Location/Qualifiers
1..56
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="SALK_096733"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna_protocols.html"
BASE COUNT      19 a      5 c      12 g      20 t
ORIGIN
Query Match      100.0%; Score 10; DB 28; Length 56;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
        |||||
Db      53 CTTCTCTTTT 44

RESULT 29
BH222307
LOCUS

```

```

DEFINITION kx19c11.v3 Parastrongyloides trichosuri FL pAMP1 v1 Chiapelli
ACCESSION MCarter Parastrongyloides trichosuri cDNA 5', mRNA sequence.
VERSION BI322307
KEYWORDS BI322307.1 GI:15001493
SOURCE Parastrongyloides trichosuri
ORGANISM Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida;
          Panagrolaimoidea; Strongyloidea; Parastrongyloides.
REFERENCE 1 (bases 1 to 57)
AUTHORS McCarter, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J., Wylie, T.,
          Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y.,
          Gibbons, M., Ritter, B., Bennett, J., Franklin, C., Tsagaris, V., R.,
          Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Steptoe
          M., Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S.,
          Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and
          Wilson, R.
TITLE The Washington Univ. Nematode EST Project, 1999
JOURNAL Unpublished
COMMENT Contact: McCarter JP
          The Washington Univ. Nematode EST Project, 1999
          Washington University School of Medicine
          4444 Forest Park Parkway, Box 8501, St. Louis, MO 63103, USA
          Tel: 314 286 1800
          Fax: 314 286 1810
          Email: est@watson.wustl.edu
          The library was constructed by Brandi Chiapelli and Dr. James
          McCarter (bchiapelli@watson.wustl.edu & jmcarter@watson.wustl.edu) at
          Washington University, St. Louis. DNA Sequencing by: Washington
          University Genome Sequencing Center St. Louis.
          Putative full length read
          The vector to vector length is 58
          Seq primer: -40RP from Gibco.
FEATURES
    source
        1..57
        /organism="Parastrongyloides trichosuri"
        /mol_type="mRNA"
        /db_xref="taxon:131310"
        /dev_stage="Free Living"
        /lab_host="DH10B"
        /clone_lib="Parastrongyloides trichosuri FL pAMP1 v1
        Chiapelli McCarter"
        /notes="Vector: pAMP1 (Gibco); Site 1: NotI; Site 2: SalI;
        The library was constructed by Brandi Chiapelli and Dr.
        James McCarter at Washington University, St. Louis. The
        cDNA was made by using Dynabead oligo-dT priming (Dyna).
        PCR based library using a modified protocol from the
        SMART PCR cDNA Synthesis Kit from Clontech. Directionally
        cloned into the UDG sites of pAMP1. Nematodes were
        provided by Dr. Warwick Grant of AgResearch, New Zealand
        (warwick.grant@agresearch.co.nz)."
BASE COUNT 15 a 6 c 4 g 32 t
ORIGIN
Query Match 100.0%; Score 10; DB 12; Length 57;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTTT 10
    |||||
Db 16 CTCTCTCTTTT 25

RESULT 30
CB917098 57 bp mRNA linear EST 25-APR-2003
LOCUS VVD118G10.371857 An expressed sequence tag database for abiotic
DEFINITION stressed berries of Vitis vinifera var. Chardonnay Vitis vinifera
          cDNA clone VVD118G10 5, mRNA sequence.
ACCESSION CB917098
VERSION CB917098.1 GI:30131759
KEYWORDS EST.
SOURCE Vitis vinifera

Query Match 100.0%; Score 10; DB 14; Length 57;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTTT 10
    |||||
Db 12 CTCTCTCTTTT 3

RESULT 31
AI343303/c 58 bp mRNA linear EST 08-APR-1999
LOCUS tb92G09.x1 NCI CGAP Lu25 Homo sapiens cDNA clone IMAGE:2061856 3'
DEFINITION similar to TR:Q33578 Q33578 KINETOPLAST CR5 1, mRNA sequence.
ACCESSION AI343303
VERSION AI343303.1 GI:4080509
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 58)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
          Tumor Gene Index
JOURNAL Unpublished
COMMENT Contact: Robert Strausberg, Ph.D.
          Email: cgapbs-@email.nih.gov
          cDNA Library Preparation: David B. Krizman, Ph.D.
          cDNA Library Arrayed by: I.M.A.G.E. Consortium, LLNL
          DNA Sequencing by: Washington University Genome Sequencing Center
          Clone distribution: NCI-CGAP clone distribution information can be
          found through the I.M.A.G.E. Consortium/LLNL at:
          www-bio.llnl.gov/bbrp/image/image.html
          Trace considered overall poor quality

```

```

ORGANISM Vitis vinifera
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; Vitaceae; Vitis.
REFERENCE 1 (bases 1 to 57)
AUTHORS Cushman, J.C.
TITLE An expressed sequence tag database for abiotic stressed berries of
          Vitis vinifera var. Chardonnay
JOURNAL Unpublished
COMMENT Contact: Cushman JC
          Department of Biochemistry
          University of Nevada
          MS200, Reno, NV 89557-0014, USA
          Tel: 775-784-1918
          Fax: 775-784-1650
          Email: jcushman@unr.edu
          PCR Primers
          FORWARD: T3 20mer
          BACKWARD: T7 21mer (backward)
          Plate: 118 row: G column: 10
          Seq primer: T3 20mer
          High quality sequence stop: 57.
          Location/Qualifiers
              1..57
              /organism="Vitis vinifera"
              /mol_type="mRNA"
              /db_xref="taxon:29760"
              /clone="VVD118G10"
              /tissue_type="berries"
              /dev_stage="mixed; 8, 9, 11, 13, 15, 16 weeks dar"
              /clone_lib="An expressed sequence tag database for abiotic
              stressed berries of Vitis vinifera var. Chardonnay"
              /notes="Vector: Lambda Uni-Zap XR, Bluescript SK-; Site_1:
              EcoRI; Site_2: XhoI"
BASE COUNT 29 a 1 c 14 g 13 t
ORIGIN
Query Match 100.0%; Score 10; DB 14; Length 57;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTTT 10
    |||||
Db 12 CTCTCTCTTTT 3

RESULT 31
AI343303/c 58 bp mRNA linear EST 08-APR-1999
LOCUS tb92G09.x1 NCI CGAP Lu25 Homo sapiens cDNA clone IMAGE:2061856 3'
DEFINITION similar to TR:Q33578 Q33578 KINETOPLAST CR5 1, mRNA sequence.
ACCESSION AI343303
VERSION AI343303.1 GI:4080509
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 58)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
          Tumor Gene Index
JOURNAL Unpublished
COMMENT Contact: Robert Strausberg, Ph.D.
          Email: cgapbs-@email.nih.gov
          cDNA Library Preparation: David B. Krizman, Ph.D.
          cDNA Library Arrayed by: I.M.A.G.E. Consortium, LLNL
          DNA Sequencing by: Washington University Genome Sequencing Center
          Clone distribution: NCI-CGAP clone distribution information can be
          found through the I.M.A.G.E. Consortium/LLNL at:
          www-bio.llnl.gov/bbrp/image/image.html
          Trace considered overall poor quality

```

Seq primer: -40UP from Gibco
High quality sequence stop: 1.

FEATURES

source
1...58
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2061856"
/tissue_type="bronchioalveolar carcinoma"
/dev_stage="adult"
/lab_host="DH10B"
/clone_lib="NCI-GAP Lu25"
/note="Organ: lung; Vector: pAMP1; mRNA made from lung carcinoma tissue, cDNA made by oligo-dr priming. Directionally cloned. Size-selected on agarose gel, average insert size 500 bp. Primary library, non-amplified." 34 a 15 c 8 g 1 t

BASE COUNT
ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 58;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 47 CTTCTCTTTT 38

RESULT 32
LOCUS BQ595228/c
DEFINITION E012710-024-023-F23-SP6 MP1Z-ADIS-024-developing root Beta vulgaris cDNA clone 024-023-F23 5-PRIME, mRNA sequence.
ACCESSION BQ595228
VERSION BQ595228.1 GI:26124811
KEYWORDS EST.
SOURCE Beta vulgaris
ORGANISM Beta vulgaris
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Caryophyllales; Amaranthaceae; Beta.
REFERENCE 1 (bases 1 to 58)
AUTHORS Herwig,R., Schulz,B., Weishaar,B., Hennig,S., Steinfath,M., Drungowski,M., Stahl,D., Wruck,W., Menze,A., O'Brien,J., Lehrach,H. and Radelof,U.
TITLE Construction of a 'unigene' cDNA clone set by oligonucleotide fingerprinting allows access to 25 000 potential sugar beet genes
JOURNAL Plant J. 32 (5), 845-857 (2002)
COMMENT Contact: Weishaar B
ADIS DNA core facility at MP1Z
Max-Planck-Institute for Plant Breeding Research
Carl-von-Linne Weg 10, 50829 Koeln, Germany
Fax: 00492215062851
Email: weissshaar@mp1z-koeln.mpg.de
Insert Length: 58 Std Error: 0.00
Plate: 23 row: F column: 23
Seq primer: SP6; CATACGATTAGTGACACTATAG.

FEATURES

source
1...58
/organism="Beta vulgaris"
/mol_type="mRNA"
/cultivar="KWS2320 (double haploid, monogerm breeding line)"
/db_xref="GABI:191621"
/db_xref="taxon:161934"
/clone="024-023-F23"
/tissue_type="developing root"
/lab_host="EMDH10B"
/clone_lib="MP1Z-ADIS-024-developing root"
/note="Vector: pCMVSPOR16; Site 1: SalI; Site 2: NotI; cDNA library from sugar beet, library provided by KWS Kleinwanzlebener Saatzzucht AG Binbeck, Germany, contact:

b.schulz@kws.de; cloning sites SalI-NotI, primer sites and orientation:
SP6-Sali-CCACGCGTCCG-5prime-cDNA-polyA-CC-NotI-T7; Note: Sequencing granted in the context of the GABI-beet project, local PI: Dr. Katharina Schneider, coordinator: Prof. Christian Jung; Sequence submission managed by RZPD/GABI-Primary database: <http://gabi.rzpd.de>

BASE COUNT 32 a 8 c 17 g 1 t
ORIGIN

Query Match 100.0%; Score 10; DB 13; Length 58;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||

Db 54 CTTCTCTTTT 45

RESULT 33

LOCUS AU077187
DEFINITION AU077187 Sugano cDNA library Homo sapiens cDNA clone Zrv61646 similar to 5'-end region of Mouse mRNA for proteasome Z subunit, mRNA sequence.
ACCESSION AU077187
VERSION AU077187.1 GI:7439801
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 59)
AUTHORS Suzuki,Y., Ishihara,D., Sasaki,M., Nakagawa,H., Hata,H., Tsunoda,T., Watanabe,M., Komatsu,T., Ota,T., Isogai,T., Suyama,A. and Sugano,S.
TITLE Statistical analysis of the 5' untranslated region of human mRNA using 'Oligo-Capped' cDNA libraries
JOURNAL Genomics 64 (3), 286-297 (2000)
MEDLINE 20221373
PUBMED 10756096
COMMENT Contact: Yutaka Suzuki
Department of Virology
Institute of Medical Science, University of Tokyo
4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
Email: ysuzuki@ims.u-tokyo.ac.jp
Suzuki,Y., Yoshitomo-Nakagawa,K., Maruyama,K., Suyama,A. and Sugano,S. Construction and characterization of a full length-enriched and a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997)
This clone was obtained from a '5'-end-enriched' cDNA library constructed by 'Oligo-Capping' method. The coding region starts from the 50 bp upstream to the 3'-end.

FEATURES

source
1...59
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="Zrv61646"
/clone_lib="Sugano cDNA library"
BASE COUNT 9 a 12 c 19 g 19 t
ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 59;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||

Db 48 CTTCTCTTTT 57

RESULT 34

AZ774319

LOCUS
DEFINITION AZ774339 59 bp DNA linear GSS 16-FEB-2001
 2M003005R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC2M003005 R, genomic survey sequence.
ACCESSION AZ774339
VERSION GSS.
KEYWORDS AZ774339.1 GI:12899665
SOURCE
ORGANISM Mus musculus (house mouse)
REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 59)
 Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,
 Islan, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly,
 M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niederhausern, A.
 and Wright, D., Weiss, R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
JOURNAL Unpublished
COMMENT Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0003 row: O column: 05
 Seq primer: CACACAGAAACAGCTATGACC
 Class: plasmid ends
 High quality sequence stop: 59.
FEATURES
 source
 Location/Qualifiers
 1..59
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC2M003005"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /notes="Vector: PWD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adaptor DNA was purified and size-selected for a 9.5 to
 10.5 Kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pWB42 (GI|4732114|gb|AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."
 12 a 7 c 12 g 28 t

Query Match 100.0%; Score 10; DB 28; Length 59;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
 |||||
Db 19 CTTCTCTTTT 29

RESULT 35

LOCUS
DEFINITION BZ354535 59 bp DNA linear GSS 14-NOV-2002
 SALK_125268.25.05.x Arabidopsis thaliana TDNA insertion lines
 Arabidopsis thaliana genomic clone SALK_125268.25.05.x, genomic
 survey sequence.
ACCESSION BZ354535
VERSION BZ354535
KEYWORDS GSS.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
 ; eurosids II; Brassicales; Brassicaceae; Arabidopsids.
 1 (bases 1 to 59)
 Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R., Gadrinab,
 C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prednis, L., Shinn, P.,
 Zimmerman, J., and Ecker, J.R.
TITLE A Sequence-Indexed Library of Insertion Mutations in the
 Arabidopsis Genome
JOURNAL Unpublished
COMMENT Contact: Joseph R. Ecker
 Salk Institute Genomic Analysis Laboratory (SIGnAL)
 The Salk Institute for Biological Studies
 10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
 Tel: 858 453 4100 x1752
 Fax: 858 558 6379
 Email: ecker@salk.edu
 This is single pass sequence recovered from the left border of
 TDNA.
Class: TDNA tagged.

FEATURES
 source
 Location/Qualifiers
 1..59
 /organism="Arabidopsis thaliana"
 /mol_type="genomic DNA"
 /strain="Columbia 0"
 /db_xref="taxon:3702"
 /clone="SALK_125268.25.05.x"
 /clone_lib="Arabidopsis thaliana TDNA insertion lines"
 /notes="PCR was performed on Arabidopsis thaliana lines
 each of which contains one or more TDNA insertion
 elements. The resultant fragment for each line was
 directly sequenced to determine the genomic sequence at
 the site of insertion. Details of the protocols used can
 be found at http://signal.salk.edu/tdna_protocols.html"
 7 a 16 c 14 g 22 t

Query Match 100.0%; Score 10; DB 29; Length 59;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
 |||||
Db 1 CTTCTCTTTT 10

RESULT 36
AL762585
LOCUS AL762585 59 bp DNA linear GSS 19-JUN-2002
 Arabidopsis thaliana T-DNA flanking sequence GK-026B10-013759,
 genomic survey sequence.
ACCESSION AL762585
VERSION AL762585.1 GI:21506841
KEYWORDS GSS.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
REFERENCE Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
 rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsids.
 1
 Strizhov, N., Li, Y., Rosso, M., Viehoveer, P., Dekker, K., Siedler, H.
 and Weisshaar, B.
TITLE A pipeline for automated high-throughput generation of FSTs

(flanking sequence tags) from Arabidopsis thaliana T-DNA

transformed lines
Unpublished

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Rosso, M., Strizhov, N., Li, Y., Reiss, B., Dekker, K. and Weishaar, B.
A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat)
for flanking sequence tag based reverse genetics
Unpublished
3 (bases 1 to 59)
Li, Y., Strizhov, N., Rosso, M. and Weishaar, B.
Direct Submission
Submitted (17-JUN-2002) Weishaar B., Max-Planck-Institut fuer
Zuechtungsforchung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany
This sequence is recovered from the left border of the T-DNA. It
indicates an insertion within the locus defined by clone F23B13.
The sequences are generated at the MPI for Plant Breeding Research
in the context of the GABI-Kat project. GABI-Kat is part of the
German Plant Genomics program designated 'GABI'. Information on
line availability can be found at:
<http://www.mpiz-koeln.mpg.de/GABI-Kat/>.

FEATURES

source

Location/Qualifiers

1..59

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Columbia 0"

/db_xref="taxon:3702"

/clone="GK-026B10-013759"

/note="PCR was performed on DNA from Arabidopsis thaliana
plants (T1) which were transformed with the T-DNA from
vector PAC106. The lines contain one or more T-DNA
insertions. The DNA fragment(s) resulting from the PCR
were directly sequenced to determine the genomic sequence
flanking the insertion. Sequences displaying significant
similarity to the A. thaliana nuclear genome sequence were
processed for submission. T-DNA derived sequences were
removed"

BASE COUNT 9 a 11 c 12 g 27 t

ORIGIN

Query Match

Best Local Similarity 100.0%; Score 10; DB 29; Length 59;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

|||||

33 CTTCTCTTTT 42

RESULT 37

AU258677

LOCUS

DEFINITION

AU258677 3'-directed mouse cDNA library Mus musculus cDNA clone

BED0013467 3', mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

63 bp mRNA linear EST 25-APR-2002
Mus musculus (house mouse)
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 63)
Kato, K. and Matoba, R.
Generation of expressed sequence tags from mouse brain
Unpublished
Contact: Kikuya Kato
Graduate School of Biological Sciences
Nara Institute of Science and Technology
8916-5 Takayama, Ikoma, Nara 630-0101, Japan
Tel: 81-743-72-5581
Fax: 81-743-72-5589
Email: kkato@bs.nara.ac.jp,
URL: <http://love2.aist-nara.ac.jp/BED/index.html>.

FEATURES

source

Location/Qualifiers

1..63

/organism="Mus musculus"

/mol_type="mRNA"

/db_xref="taxon:10090"

/clone="BED0013467"

/tissue_type="brain"

/clone_lib="3'-directed mouse cDNA library"

BASE COUNT 16 a 11 c 11 g 24 t

ORIGIN

Query Match

Best Local Similarity 100.0%; Score 10; DB 9; Length 63;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

|||||

15 CTTCTCTTTT 24

RESULT 38

AW874904/c

LOCUS

DEFINITION

AW874904 64 bp mRNA linear EST 22-MAY-2000

SWYACAL04A01SK Brugia malayi young adult cDNA (SAW99MLW-BmYA)

Brugia malayi cDNA clone SWYACAL04A01 5', mRNA sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

AW874904 64 bp mRNA linear EST 22-MAY-2000
SWYACAL04A01SK Brugia malayi young adult cDNA (SAW99MLW-BmYA)
Brugia malayi cDNA clone SWYACAL04A01 5', mRNA sequence.
AW874904 1 GI:8012605
EST.
Brugia malayi
Brugia malayi
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Brugia.
1 (bases 1 to 64)
Williams, S.A.
Genes expressed in young adult of Brugia malayi
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers

FEATURES

source

1..64

/organism="Brugia malayi"

/mol_type="mRNA"

/db_xref="taxon:6279"

/clone="SWYACAL04A01"

/dev_stage="young adult"

/lab_host="XLI-Blue MRF"

/clone_lib="Brugia malayi young adult cDNA (SAW99MLW-BmYA)"

/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Lymphatic filarial nematode parasite of humans.
mRNA was prepared from young adult worms isolated from the
peritoneal cavity of jirds and converted to
double-stranded cDNA using reverse transcriptase and
oligo(dT) followed by RNase H and DNA pol I. The library
has 6.5 x 104 independent recombinants and the average
insert size is approx. 800bp. The library was constructed
by Michelle Lizotte-Waniewski. The library is available
from Dr. S.A. Williams, email: genome@smith.edu."

25 a 10 c 18 g 11 t

BASE COUNT

ORIGIN

Query Match

Best Local Similarity 100.0%; Score 10; DB 10; Length 64;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10


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/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVL2CAS08G03"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library is constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT      25 a      9 c      19 g      11 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 14 CTTCTCTTTT 5

RESULT 42
BE239293/c
LOCUS
DEFINITION
SWOVL2CAS08H03SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
ONCHOCERCIDAE; Onchocerca.
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Williams, S.A.
Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. 64
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/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVL2CAS08H03"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library is constructed by Michelle Lizotte-Waniewski. The library is

```

```

available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT      29 a      11 c      15 g      9 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 47 CTTCTCTTTT 38

RESULT 43
BE636299/c
LOCUS
DEFINITION
SMOVL2CAS13H10SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
ONCHOCERCIDAE; Onchocerca.
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Williams, S.A.
Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. 64
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/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SMOVL2CAS13H10"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library is constructed by Michelle Lizotte-Waniewski. The library is

```

```

available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT      29 a      11 c      14 g      10 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
    |||||
Db 38 CTTCTCTTTT 29

RESULT 44
BE636308/c
LOCUS
DEFINITION
SMOVL2CAS13H10SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
ONCHOCERCIDAE; Onchocerca.
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Williams, S.A.
Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. 64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SMOVL2CAS13H10"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library is constructed by Michelle Lizotte-Waniewski. The library is

```

DEFINITION SMOVL2CASI4A09SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
Onchocerca volvulus cDNA clone SMOVL2CASI4A09 5', mRNA sequence.

ACCESSION BE636308
VERSION BE636308.1 GI:9919515
KEYWORDS EST.

SOURCE ORGANISM

Onchocerca volvulus

Onchocerca volvulus

Onchocercidae; Onchocerca.
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;

1 (bases 1 to 64)

Genes expressed in L2 larvae of Onchocerca volvulus

Unpublished

Contact: Steven A. Williams

Molecular Parasitology

Smith College Department of Biological Sciences

Department of Biological Sciences, Clark Science Center, Smith

College, Northampton, MA, 01063, USA

Tel: 4135853826

Fax: 4135853786

Email: genome@smith.edu

Seq primer: pBluescript SK.

Location/Qualifiers

FEATURES

source

1..64

/organism="Onchocerca volvulus"

/mol_type="mRNA"

/db_xref="taxon:6282"

/clone="SMOVL2CASI4A09"

/dev_stage="L2"

/lab_host="XLI-Blue MRF"

/clone_lib="Onchocerca volvulus L2 larvae cDNA

(SAW98MLW-OvL2)"

/notes="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:

Xho I; Filarial nematode parasite of humans. mRNA was

prepared from approximately 9,000 L2s isolated from

infected mosquitoes from Kumba, Cameroon and converted to

double-stranded cDNA using reverse transcriptase and

oligo(dT) followed by RNase H and DNA pol I. The library

has 7.3 x 10⁶ independent recombinants and the average

insert size is approximately 1kb. The library was

constructed by Michelle Lizotte-Waniewski. The library is

available from Dr.S.A.Williams, email: genome@smith.edu."

23 a 9 c 19 g 13 t

BASE COUNT

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTT 10

Db 11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

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11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

11 CTCTCTCTTT 2

Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA

Tel: 4135853826

Fax: 4135853786

Email: genome@smith.edu

Seq primer: pBluescript SK.

Location/Qualifiers

FEATURES

source

1..64

/organism="Onchocerca volvulus"

/mol_type="mRNA"

/db_xref="taxon:6282"

/clone="SMOVL2CASI4C02"

/dev_stage="L2"

/lab_host="XLI-Blue MRF"

/clone_lib="Onchocerca volvulus L2 larvae cDNA

(SAW98MLW-OvL2)"

/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:

Xho I; Filarial nematode parasite of humans. mRNA was

prepared from approximately 9,000 L2s isolated from

infected mosquitoes from Kumba, Cameroon and converted to

double-stranded cDNA using reverse transcriptase and

oligo(dT) followed by RNase H and DNA pol I. The library

has 7.3 x 10⁶ independent recombinants and the average

insert size is approximately 1kb. The library was

constructed by Michelle Lizotte-Waniewski. The library is

available from Dr.S.A.Williams, email: genome@smith.edu."

26 a 8 c 19 g 11 t

BASE COUNT

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTT 10

Db 16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

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16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

16 CTCTCTCTTT 7

RESULT 46

BE636344/c

LOCUS

DEFINITION

Onchocerca volvulus cDNA clone SMOVL2CASI4E08 5', mRNA sequence.

ACCESSION

BE636344

VERSION

BE636344.1 GI:9919551

KEYWORDS

EST.

SOURCE

Onchocerca volvulus

Onchocerca volvulus

Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;

Onchocercidae; Onchocerca.

1 (bases 1 to 64)

Williams,S.A.

Genes expressed in L2 larvae of Onchocerca volvulus

Unpublished

Contact: Steven A. Williams

Molecular Parasitology

Smith College Department of Biological Sciences

Department of Biological Sciences, Clark Science Center, Smith

College, Northampton, MA, 01063, USA

Tel: 4135853826

Fax: 4135853786

Email: genome@smith.edu

Seq primer: pBluescript SK.

Location/Qualifiers

FEATURES

source

1..64

/organism="Onchocerca volvulus"

/mol_type="mRNA"

/db_xref="taxon:6282"

/clone="SMOVL2CASI4E08"

/dev_stage="L2"

/lab_host="XLI-Blue MRF"

/clone_lib="Onchocerca volvulus L2 larvae cDNA

(SAW98MLW-OvL2)"

/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT 30 a 10 c 14 g 10 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 38 CTTCTCTTTT 29

RESULT 47
BE636363/c
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

BE636363 64 bp mRNA linear EST 25-AUG-2000
SWOVL2CAS14G07SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
Onchocerca volvulus cDNA clone SWOVL2CAS14G07 5', mRNA sequence.

BE636363
EST
Onchocerca volvulus
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
1 (bases 1 to 64)
Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pbluescript SK.
Location/Qualifiers
1..64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVL2CAS14G07"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)"

/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT 27 a 8 c 18 g 11 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 11 CTTCTCTTTT 2

RESULT 49
BE636392/c
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

BE636392 64 bp mRNA linear EST 25-AUG-2000
SWOVL2CAS14H03SK Onchocerca volvulus adult male cDNA (SAW98MLW-OvL2)
Onchocerca volvulus cDNA clone SWOVL2CAS14H03 5', mRNA sequence.

BE636392
EST
Onchocerca volvulus
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
1 (bases 1 to 64)
Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pbluescript SK.
Location/Qualifiers
1..64
/organism="Onchocerca volvulus"
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/db_xref="taxon:6282"
/clone="SWOVL2CAS14G07"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)"

/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT 23 a 9 c 19 g 13 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 11 CTTCTCTTTT 2

RESULT 49
BE636392/c
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

BE636392 64 bp mRNA linear EST 25-AUG-2000
SWOVL2CAS14H03SK Onchocerca volvulus adult male cDNA (SAW98MLW-OvL2)
Onchocerca volvulus cDNA clone SWOVL2CAS14H03 5', mRNA sequence.

BE636392
EST
Onchocerca volvulus
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
1 (bases 1 to 64)
Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pbluescript SK.
Location/Qualifiers
1..64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVL2CAS14H03"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)"

Qy 1 CTTCTCTTTT 10
|||||
Db 18 CTTCTCTTTT 9

RESULT 48
BE636369/c

LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

BE636369 64 bp mRNA linear EST 25-AUG-2000
SWOVL2CAS14H03SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
Onchocerca volvulus cDNA clone SWOVL2CAS14H03 5', mRNA sequence.

BE636369
EST
Onchocerca volvulus
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
1 (bases 1 to 64)
Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pbluescript SK.
Location/Qualifiers
1..64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVL2CAS14H03"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)"

/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT 23 a 9 c 19 g 13 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 11 CTTCTCTTTT 2

RESULT 49
BE636392/c
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

BE636392 64 bp mRNA linear EST 25-AUG-2000
SWOVL2CAS14H03SK Onchocerca volvulus adult male cDNA (SAW98MLW-OvL2)
Onchocerca volvulus cDNA clone SWOVL2CAS14H03 5', mRNA sequence.

BE636392
EST
Onchocerca volvulus
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
1 (bases 1 to 64)
Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pbluescript SK.
Location/Qualifiers
1..64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVL2CAS14H03"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)"

/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10E4 independent recombinants and the average insert size is approximately 1kb. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT 23 a 9 c 19 g 13 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 11 CTTCTCTTTT 2

RESULT 49
BE636392/c
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

BE636392 64 bp mRNA linear EST 25-AUG-2000
SWOVL2CAS14H03SK Onchocerca volvulus adult male cDNA (SAW98MLW-OvL2)
Onchocerca volvulus cDNA clone SWOVL2CAS14H03 5', mRNA sequence.

BE636392
EST
Onchocerca volvulus
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
1 (bases 1 to 64)
Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
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Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pbluescript SK.
Location/Qualifiers
1..64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVL2CAS14H03"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)"

Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.

1 (bases 1 to 64)
Lizotte-Waniewski, M. and Williams, S.A.
Genes expressed in adult male stage of *Onchocerca volvulus*
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu

Seq primer: pBluescript SK.

Location/Qualifiers
1. .64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVAMCAQ09D08"
/sex="male"
/dev_stage="adult"
/lab_host="XLI-Blue MRP"
/clone_lib="Onchocerca volvulus adult male cDNA
(SAW98MLW-OvAM)"
/notes="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. Six adult
male worms of *Onchocerca volvulus* were isolated from
consenting patients and quick frozen. Adult male mRNA was
converted to double-stranded cDNA using reverse
transcriptase and oligo(dT) followed by RNase H and DNA
pol I. The library has 2 x 10E5 independent recombinants
and the average insert size is ~1100bp. The library was
constructed by Michelle Lizotte-Waniewski with worms
provided by Dr. Sara Lustigman. The library is available
from Dr. Steven A. Williams, email: genome@smith.edu."

BASE COUNT 30 a 11 c 13 g 10 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
|||||
Db 54 CTTCTCTTTT 45

RESULT 50
BE636430/c 64 bp mRNA linear EST 25-AUG-2000
LOCUS
DEFINITION SWOVLCAS13C02SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)

ACCESSION BE636430
VERSION BE636430.1 GI:9919457
KEYWORDS EST.
SOURCE Onchocerca volvulus
ORGANISM Onchocercidae; Onchocerca.

REFERENCE
AUTHORS Smith College Department of Biological Sciences
TITLE Department of Biological Sciences, Clark Science Center, Smith
JOURNAL College, Northampton, MA, 01063, USA
COMMENT Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu

Seq primer: pBluescript SK.
Location/Qualifiers
1. .64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVLCAS13C02SK"
/dev_stage="L2"
/lab_host="XLI-Blue MRP"
/clone_lib="Onchocerca volvulus L2 larvae cDNA
(SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. mRNA was
prepared from approximately 9,000 L2s isolated from
infected mosquitoes from Kumba, Cameroon and converted to
double-stranded cDNA using reverse transcriptase and

BASE COUNT 30 a 11 c 13 g 10 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
|||||
Db 54 CTTCTCTTTT 45

RESULT 50
BE636430/c 64 bp mRNA linear EST 25-AUG-2000
LOCUS
DEFINITION SWOVLCAS13C02SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)

ACCESSION BE636430
VERSION BE636430.1 GI:9919457
KEYWORDS EST.
SOURCE Onchocerca volvulus
ORGANISM Onchocercidae; Onchocerca.

FEATURES

source

Seq primer: pBluescript SK.
Location/Qualifiers
1. .64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVLCAS13C02"
/dev_stage="L2"
/lab_host="XLI-Blue MRP"
/clone_lib="Onchocerca volvulus L2 larvae cDNA
(SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. mRNA was
prepared from approximately 9,000 L2s isolated from
infected mosquitoes from Kumba, Cameroon and converted to
double-stranded cDNA using reverse transcriptase and
oligo(dT) followed by RNase H and DNA pol I. The library
has 7.3 x 10E4 independent recombinants and the average
insert size is approximately 1kb. The library was
constructed by Michelle Lizotte-Waniewski. The library is
available from Dr. S.A. Williams, email: genome@smith.edu."

BASE COUNT 28 a 9 c 15 g 12 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
|||||
Db 33 CTTCTCTTTT 24

RESULT 51

BE636451/c

LOCUS

DEFINITION BE636451

ACCESSION BE636451

VERSION BE636451.1 GI:9919478

KEYWORDS EST.

SOURCE Onchocerca volvulus

ORGANISM Onchocercidae; Onchocerca.

REFERENCE 1 (bases 1 to 64)

AUTHORS Williams, S.A.

TITLE Genes expressed in L2 larvae of *Onchocerca volvulus*

JOURNAL Unpublished

COMMENT Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu

FEATURES

source

Seq primer: pBluescript SK.
Location/Qualifiers
1. .64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVLCAS13B09"
/dev_stage="L2"
/lab_host="XLI-Blue MRP"
/clone_lib="Onchocerca volvulus L2 larvae cDNA
(SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. mRNA was
prepared from approximately 9,000 L2s isolated from
infected mosquitoes from Kumba, Cameroon and converted to
double-stranded cDNA using reverse transcriptase and

BASE COUNT 28 a 9 c 15 g 12 t

ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
|||||
Db 33 CTTCTCTTTT 24

RESULT 51

BE636451/c

LOCUS

DEFINITION BE636451

ACCESSION BE636451

VERSION BE636451.1 GI:9919478

KEYWORDS EST.

SOURCE Onchocerca volvulus

ORGANISM Onchocercidae; Onchocerca.

REFERENCE 1 (bases 1 to 64)

AUTHORS Williams, S.A.

TITLE Genes expressed in L2 larvae of *Onchocerca volvulus*

JOURNAL Unpublished

COMMENT Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu

oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10⁵ independent recombinants and the average insert size is approximately 1kb. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT
ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 16 CTTCTCTTTT 7

RESULT 52
BE636456/c

LOCUS SWOV12CAS13F038K Onchocerca volvulus L2 larvae cDNA (SAW98MLW-Ovl2)
DEFINITION Onchocerca volvulus cDNA clone SWOV12CAS13F03 5', mRNA sequence.

ACCESSION BE636456
VERSION BE636456.1 GI:9919483
KEYWORDS EST.

SOURCE ORGANISM

Onchocerca volvulus
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.

1 (bases 1 to 64)
Williams, S.A.

Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished

Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.

FEATURES
source

1..64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOV12CAS13F03"
/dev_stage="L2"
/lab_host="XLI-Blue MRP"
/clone_lib="Onchocerca volvulus L2 larvae cDNA (SAW98MLW-Ovl2)"

/note="vector: Lambda Uni-ZAP XR; Site_1: Eco RI; Site_2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10⁵ independent recombinants and the average insert size is approximately 1kb. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT
ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 41 CTTCTCTTTT 32

RESULT 53
BE636464/c

LOCUS SWOV12CAS13G02SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-Ovl2)
DEFINITION Onchocerca volvulus cDNA clone SWOV12CAS13G02 5', mRNA sequence.

ACCESSION BE636464
VERSION BE636464.1 GI:9919491
KEYWORDS EST.

SOURCE ORGANISM

Onchocerca volvulus
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.

1 (bases 1 to 64)
Williams, S.A.

Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished

Contact: Steven A. Williams
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Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.

Location/Qualifiers
1..64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOV12CAS13G02"
/dev_stage="L2"
/lab_host="XLI-Blue MRP"
/clone_lib="Onchocerca volvulus L2 larvae cDNA (SAW98MLW-Ovl2)"

/note="vector: Lambda Uni-ZAP XR; Site_1: Eco RI; Site_2: Xho I; Filarial nematode parasite of humans. mRNA was prepared from approximately 9,000 L2s isolated from infected mosquitoes from Kumba, Cameroon and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library has 7.3 x 10⁵ independent recombinants and the average insert size is approximately 1kb. The library was constructed by Michelle Lizotte-Waniewski. The library is available from Dr.S.A.Williams, email: genome@smith.edu."

BASE COUNT
ORIGIN

Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 35 CTTCTCTTTT 26

RESULT 54
BE636472/c

LOCUS SWOV12CAS13H01SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-Ovl2)
DEFINITION Onchocerca volvulus cDNA clone SWOV12CAS13H01 5', mRNA sequence.

ACCESSION BE636472
VERSION BE636472.1 GI:9919499
KEYWORDS EST.

SOURCE ORGANISM

Onchocerca volvulus
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.

1 (bases 1 to 64)
Williams, S.A.

Genes expressed in L2 larvae of Onchocerca volvulus
Unpublished

Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. .64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SMOVL2CASI3H01"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA
(SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. mRNA was
prepared from approximately 9,000 L2s isolated from
infected mosquitoes from Kumba, Cameroon and converted to
double-stranded cDNA using reverse transcriptase and
oligo(dT) followed by RNase H and DNA pol I. The library
has 7.3 x 10E4 independent recombinants and the average
insert size is approximately 1kb. The library was
constructed by Michelle Lizotte-Waniewski. The library is
available from Dr.S.A.Williams, email: genome@smith.edu."
26 a 7 c 19 g 12 t

BASE COUNT
ORIGIN
Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 21 CTTCTCTTTT 12

RESULT 55
BF118488/c
LOCUS
DEFINITION
BF118488 64 bp mRNA linear EST 24-OCT-2000
SMOVL3CAN69E12SK Onchocerca volvulus infective larva cDNA
(SAW94WL-OvL3) Onchocerca volvulus cDNA clone SMOVL3CAN69E12 5',
mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
BF118488.1 GI:10992964
Onchocerca volvulus
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.
1 (bases 1 to 64)
Williams, S.A., Lu, W., Lizotte-Waniewski, M. and Laney, S.J.
Genes expressed in infective third stage larvae of Onchocerca
volvulus
1 (bases 1 to 64)
Williams, S.A., Lu, W., Lizotte-Waniewski, M. and Laney, S.J.
Genes expressed in infective third stage larvae of Onchocerca
volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. .64
/organism="Onchocerca volvulus"
/mol_type="mRNA"

Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. .64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SMOVL3CAN69E12"
/dev_stage="L3"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus molting L3 larva cDNA
(SL96MLW-Ovml3)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. Third-stage
larvae, L3, were isolated from infected black flies in
Cameroon (forest strain). The L3 were cultured in 20% FCS
in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in
culture. L3 of O. volvulus molt to fourth-stage larvae by
day 5 in culture. mRNA was isolated from approximately
6000 molting larvae (ml3). 2000 larvae from day 1, 2 or 3

BASE COUNT
ORIGIN
Query Match 100.0%; Score 10; DB 10; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 18 CTTCTCTTTT 9

RESULT 56
BI097426/c
LOCUS
DEFINITION
BI097426 64 bp mRNA linear EST 25-JUN-2001
SMOVL3CAN63F09SK Onchocerca volvulus molting L3 larva cDNA
(SL96MLW-Ovml3) Onchocerca volvulus cDNA clone SMOVL3CAN63F09 5',
mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
BI097426
BI097426.1 GI:14549083
Onchocerca volvulus
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.
1 (bases 1 to 64)
Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.
Genes expressed in molting L3 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1. .64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/strain="Kumba, Cameroons"
/db_xref="taxon:6282"
/clone="SMOVL3CAN63F09"
/dev_stage="molting L3"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus molting L3 larva cDNA
(SL96MLW-Ovml3)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. Third-stage
larvae, L3, were isolated from infected black flies in
Cameroon (forest strain). The L3 were cultured in 20% FCS
in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in
culture. L3 of O. volvulus molt to fourth-stage larvae by
day 5 in culture. mRNA was isolated from approximately
6000 molting larvae (ml3). 2000 larvae from day 1, 2 or 3

in culture, and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library was constructed in the lambda Uni-Zap XR vector and has 1 x 10E6 independent recombinants and the average insert size is ~1200 bp. The library was constructed by Sara Lustigman and Michelle Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams. The library is available from Dr. Sara Lustigman (email: slustigman@bc.org)."

BASE COUNT 30 a 11 c 13 g 10 t
 ORIGIN
 Query Match 100.0%; Score 10; DB 12; Length 64;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTT 10
 |||||
 Db 40 CTCTCTCTTT 31

RESULT 57
 BI142401/c
 LOCUS
 DEFINITION SWOV3MCAM62B04SK Onchocerca volvulus molting L3 larva cDNA
 (SL96MLW-Ovml3) Onchocerca volvulus cDNA clone SWOV3MCAM62B04 5',
 mRNA sequence.

ACCESSION BI142401.1 GI:14624111
 VERSION
 KEYWORDS
 SOURCE

ORGANISM Onchocerca volvulus
 Onchocerca volvulus
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
 Onchocercidae; Onchocerca.

REFERENCE 1 (bases 1 to 64)
 AUTHORS Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.
 TITLE Genes expressed in molting L3 larvae of Onchocerca volvulus
 JOURNAL Unpublished
 COMMENT Contact: Steven A. Williams
 Molecular Parasitology
 Smith College Department of Biological Sciences
 Department of Biological Sciences, Clark Science Center, Smith
 College, Northampton, MA, 01063, USA
 Tel: 4135853826
 Fax: 4135853786
 Email: genome@smith.edu
 Seq primer: pBluescript SK.
 Location/Qualifiers

FEATURES
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 1..64
 /organism="Onchocerca volvulus"
 /mol_type="mRNA"
 /strain="Kumba, Cameroons"
 /db_xref="taxon:6282"
 /clone="SWOV3MCAM62B04"
 /dev_stage="molting L3"
 /lab_host="XLI-Blue MRF"
 /clone_lib="Onchocerca volvulus molting L3 larva cDNA (SL96MLW-Ovml3)"
 /note="vector: Lambda Uni-Zap XR; Site_1: Eco RI; Site_2: Xho I; Filarial nematode parasite of humans. Third-stage larvae, L3, were isolated from infected black flies in Cameroon (forest strain). The L3 were cultured in 20% FCS in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in culture. L3 of O. volvulus molt to fourth-stage larvae by day 5 in culture. mRNA was isolated from approximately 6000 molting larvae (mL3), 2000 larvae from day 1, 2 or 3 in culture, and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library was constructed in the lambda Uni-Zap XR vector and has 1 x 10E6 independent recombinants and the average insert size is ~1200 bp. The library was constructed by Sara Lustigman and Michelle Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams."

The library is available from Dr. Sara Lustigman (email: slustigman@bc.org)."

BASE COUNT 27 a 10 c 15 g 12 t
 ORIGIN
 Query Match 100.0%; Score 10; DB 12; Length 64;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTCTTT 10
 |||||
 Db 35 CTCTCTCTTT 26

RESULT 58
 BI142408/c
 LOCUS
 DEFINITION SWOV3MCAM62B12SK Onchocerca volvulus molting L3 larva cDNA
 (SL96MLW-Ovml3) Onchocerca volvulus cDNA clone SWOV3MCAM62B12 5',
 mRNA sequence.

ACCESSION BI142408.1 GI:14624118
 VERSION
 KEYWORDS
 SOURCE

ORGANISM Onchocerca volvulus
 Onchocerca volvulus
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
 Onchocercidae; Onchocerca.

REFERENCE 1 (bases 1 to 64)
 AUTHORS Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.
 TITLE Genes expressed in molting L3 larvae of Onchocerca volvulus
 JOURNAL Unpublished
 COMMENT Contact: Steven A. Williams
 Molecular Parasitology
 Smith College Department of Biological Sciences
 Department of Biological Sciences, Clark Science Center, Smith
 College, Northampton, MA, 01063, USA
 Tel: 4135853826
 Fax: 4135853786
 Email: genome@smith.edu
 Seq primer: pBluescript SK.
 Location/Qualifiers

FEATURES
 source
 1..64
 /organism="Onchocerca volvulus"
 /mol_type="mRNA"
 /strain="Kumba, Cameroons"
 /db_xref="taxon:6282"
 /clone="SWOV3MCAM62B12"
 /dev_stage="molting L3"
 /lab_host="XLI-Blue MRF"
 /clone_lib="Onchocerca volvulus molting L3 larva cDNA (SL96MLW-Ovml3)"
 /note="vector: Lambda Uni-Zap XR; Site_1: Eco RI; Site_2: Xho I; Filarial nematode parasite of humans. Third-stage larvae, L3, were isolated from infected black flies in Cameroon (forest strain). The L3 were cultured in 20% FCS in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in culture. L3 of O. volvulus molt to fourth-stage larvae by day 5 in culture. mRNA was isolated from approximately 6000 molting larvae (mL3), 2000 larvae from day 1, 2 or 3 in culture, and converted to double-stranded cDNA using reverse transcriptase and oligo(dT) followed by RNase H and DNA pol I. The library was constructed in the lambda Uni-Zap XR vector and has 1 x 10E6 independent recombinants and the average insert size is ~1200 bp. The library was constructed by Sara Lustigman and Michelle Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams. The library is available from Dr. Sara Lustigman (email: slustigman@bc.org)."

BASE COUNT 28 a 8 c 17 g 11 t
 ORIGIN

Query Match 100.0%; Score 10; DB 12; Length 64;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 30 CTTCTCTTTT 21

RESULT 59
BI142422/c
LOCUS
DEFINITION
SWOV3MCAM62D04SK Onchocerca volvulus molting L3 larva cDNA
(SL96MLW-Ovml3) Onchocerca volvulus cDNA clone SWOV3MCAM62D04 5',
mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.
1 (bases 1 to 64)
Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.
Genes expressed in molting L3 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pbluescript SK.

FEATURES
Source
1..64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/strain="Kumba, Cameroons"
/db_xref="taxon:6282"
/clone="SWOV3MCAM62D04"
/dev_stage="molting L3"
/lab_host="XL1-Blue MRF"
/clone_lib="Onchocerca volvulus molting L3 larva cDNA
(SL96MLW-Ovml3)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. Third-stage
larvae, L3, were isolated from infected black flies in
Cameroon (forest strain). The L3 were cultured in 20% FCS
in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in
culture. L3 of O. volvulus molt to fourth-stage larvae by
day 5 in culture. mRNA was isolated from approximately
6000 molting larvae (mL3), 2000 larvae from day 1, 2 or 3
in culture, and converted to double-stranded cDNA using
reverse transcriptase and oligo(dT) followed by RNase H
and DNA pol I. The library was constructed in the lambda
Uni-Zap XR vector and has 1 x 10E6 independent
recombinants and the average insert size is ~1200 bp. The
library was constructed by Sara Lustigman and Michelle
Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams.
The library is available from Dr. Sara Lustigman (email:
slustig@nycbc.org)."

BASE COUNT 28 a 9 c 15 g 12 t
ORIGIN

Query Match 100.0%; Score 10; DB 12; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 34 CTTCTCTTTT 25

RESULT 60
BI142452/c
LOCUS
DEFINITION
BI142452 64 bp mRNA linear EST 05-JUL-2001
SWOV3MCAM62F12SK Onchocerca volvulus molting L3 larva cDNA
(SL96MLW-Ovml3) Onchocerca volvulus cDNA clone SWOV3MCAM62F12 5',
mRNA sequence.

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.
1 (bases 1 to 64)
Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.
Genes expressed in molting L3 larvae of Onchocerca volvulus
Unpublished
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pbluescript SK.

FEATURES
Source
1..64
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/strain="Kumba, Cameroons"
/db_xref="taxon:6282"
/clone="SWOV3MCAM62F12"
/dev_stage="molting L3"
/lab_host="XL1-Blue MRF"
/clone_lib="Onchocerca volvulus molting L3 larva cDNA
(SL96MLW-Ovml3)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. Third-stage
larvae, L3, were isolated from infected black flies in
Cameroon (forest strain). The L3 were cultured in 20% FCS
in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in
culture. L3 of O. volvulus molt to fourth-stage larvae by
day 5 in culture. mRNA was isolated from approximately
6000 molting larvae (mL3), 2000 larvae from day 1, 2 or 3
in culture, and converted to double-stranded cDNA using
reverse transcriptase and oligo(dT) followed by RNase H
and DNA pol I. The library was constructed in the lambda
Uni-Zap XR vector and has 1 x 10E6 independent
recombinants and the average insert size is ~1200 bp. The
library was constructed by Sara Lustigman and Michelle
Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams.
The library is available from Dr. Sara Lustigman (email:
slustig@nycbc.org)."

BASE COUNT 27 a 10 c 15 g 12 t
ORIGIN

Query Match 100.0%; Score 10; DB 12; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 35 CTTCTCTTTT 26

RESULT 61
AZ821452
LOCUS
DEFINITION
AZ821452 64 bp DNA linear GSS 20-FEB-2001
2M0094B14F Mouse 10kb plasmid UUGCLM library Mus musculus genomic
clone UUGC2M0094B14 F, genomic survey sequence.

ACCESSION
VERSION
AZ821452.1 GI:12991360

```

GSS.
KEYWORDS Mus musculus (house mouse)
SOURCE Mus musculus
ORGANISM Mus musculus

REFERENCE
AUTHORS Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly
M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.
and Wright,D., Weiss,R.
TITLE Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
JOURNAL Unpublished
COMMENT Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert Length: 10000 Std Error: 0.00
Plate: 0094 row: B column: 14
Seq primer: CTTTGTAAACGACGGCCAGT
Class: plasmid ends
High quality sequence stop: 64.
Location/Qualifiers
1. .64
/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUGC2M0094B14"
/sex="Male"
/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
/clone_lib="Mouse 10kb plasmid UUGC1M library"
/notes="Vector: PWP42nv; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptored DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pWD42 (gi|4732114|gb|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptored mouse DNA was annealed to
adaptored vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT 14 a 12 c 12 g 26 t
ORIGIN
Query Match 100.0%; Score 10; DB 28; Length 64;
Best Local Similarity 100.0%; Pred. NO. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 16 CTTCTCTTTT 25

RESULT 62
CB030041/c
LOCUS CB030041
DEFINITION TgESTzyl8e02.y1 TgrH Tachyzoite Norm 7 cDNA Library Toxoplasma
gondii cDNA clone TgESTzyl8e02.y1 5', mRNA sequence.
ACCESSION CB030041
CB030041 65 bp mRNA linear EST 13-JAN-2003

GSS.
KEYWORDS Mus musculus (house mouse)
SOURCE Mus musculus
ORGANISM Mus musculus

REFERENCE
AUTHORS Tang,K., Cole,R., Fogarty,S., Sibley,L.D., Ajioke,J.A., White,M.,
Clifton,S., Pape,D., Martin,J., Wylie,T., Dante,M., Marra,M.,
Hillier,L., Kucaba,T., Theising,B., Bowers,Y., Gibbons,M., Ritter
E., Bennett,J., Franklin,C., Tsagarisvili,R., Ronko,I., Kennedy
S., Maguire,L., Waterston,R. and Wilson,R.
TITLE Toxoplasma EST Project
JOURNAL Unpublished
COMMENT Contact: Clifton, S.
Toxoplasma EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: toxo@watson.wustl.edu
Contact David Sibley (toxost@borcim.wustl.edu) for further
information relating to organism, libraries, or clone availability.
Seq primer: -40RP from Gibco.
Location/Qualifiers
1. .85
/organism="Toxoplasma gondii"
/mol_type="mRNA"
/strain="RH (Type I)"
/db_xref="taxon:5811"
/clone="TgESTzyl8e02.y1"
/dev_stage="Tachyzoite"
/lab_host="DH10B (GeneHog, Invitrogen, Inc.)"
/clone_lib="TgrH Tachyzoite Norm 7 cDNA Library"
/notes="Vector: pBluscript SK-; Site 1: EcoRI; Site 2: XhoI
; Toxoplasma RH strain tachyzoites were grown in human
foreskin fibroblast cultures in vitro. The library was
originally constructed by K.L.Wan, Cambridge University.
cDNAs were synthesized from polyA RNAs by oligo d(T)
priming and directionally cloned into the EcoRI to XhoI
sites of the Lambda ZapII vector using the ZAP-cDNA
synthesis kit (Stratagene). The primary cDNA library was
mass excised as phagemid using ExAssist helper phage
(Stratagene). Phagemid DNA was extracted by
phenol-chloroform method, and hybridized against a pool of
highly abundant genes which were derived from short-cycle
PCR of the primary cDNA library. The normalized library
was electroporated into DH10B (GeneHog, Invitrogen, Inc).
WARNING: the library contains a small percentage of cDNAs
derived from the human host cells."

BASE COUNT 31 a 7 c 20 g 7 t
ORIGIN
Query Match 100.0%; Score 10; DB 14; Length 65;
Best Local Similarity 100.0%; Pred. NO. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 58 CTTCTCTTTT 49

RESULT 63
AZ514453/c
LOCUS AZ514453
DEFINITION IM0361N14F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
clone UUGC1M0361N14 F, genomic survey sequence.
ACCESSION AZ514453
VERSION AZ514453
KEYWORDS GSS.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

```

ORGANISM Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
 ; eurosids II; Brassicales; Brassicaceae; Arabidopsids.
 REFERENCE 1 (bases 1 to 65)
 AUTHORS Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab
 , C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.,
 Zimmerman,J., and Ecker,J.R.
 TITLE A Sequence-Indexed Library of Insertion Mutations in the
 Arabidopsis Genome
 JOURNAL Unpublished
 COMMENT Contact: Joseph R. Ecker
 Salk Institute Genomic Analysis Laboratory (SIGAL)
 The Salk Institute for Biological Studies
 10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
 Tel: 858 453 4100 x1752
 Fax: 858 558 6379
 Email: ecker@salk.edu
 This is single pass sequence recovered from the left border of
 TDNA. This sequence lies within 300 bases of the 5' end of
 At5g15860.
 Class: TDNA tagged.
 Location/Qualifiers
 1. .65
 /organism="Arabidopsis thaliana"
 /mol_type="genomic DNA"
 /strain="Columbia 0"
 /db_xref="taxon:3702"
 /clone="SALK_059169.41.80.x"
 /clone_lib="Arabidopsis thaliana TDNA insertion lines"
 /notes="PCR was performed on Arabidopsis thaliana lines
 each of which contains one or more TDNA insertion
 elements. The resultant fragment for each line was
 directly sequenced to determine the genomic sequence at
 the site of insertion. Details of the protocols used can
 be found at http://signal.salk.edu/tdna_protocols.html"

BASE COUNT 33 a 7 c 14 g 11 t
 ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 65;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 Db 41 CTTCTCTTTT 32

RESULT 65
 AG216203
 LOCUS AG216203
 DEFINITION Drosophila melanogaster DNA, clone NP1611-5-1, flanking P[GAWB]
 transposon insertion, genomic survey sequence.
 ACCESSION AG216203
 VERSION AG216203.1 GI:22763203
 KEYWORDS GSS.
 SOURCE Drosophila melanogaster (fruit fly)
 ORGANISM Drosophila melanogaster
 Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 Ephydroidea; Drosophilidae; Drosophila.

REFERENCE 1
 AUTHORS Hayashi,S., Ito,K., Sado,Y., Taniguchi,M., Akimoto,A., Takeuchi,H.,
 Aigaki,T., Matsuzaki,F., Nakagoshi,H., Tanimura,T., Ueda,R.,
 Uemura,T., Yoshihara,M. and Goto,S.
 TITLE GEMDB, a database compiling expression patterns and molecular
 locations of a collection of Gal4 enhancer traps
 JOURNAL Genesis (2002) In press
 REFERENCE 2 (bases 1 to 65)
 AUTHORS Hayashi,S.
 TITLE Direct Submission
 JOURNAL Submitted (27-AUG-2002) Shigeo Hayashi, RIKEN Center for
 Developmental Biology, Laboratory for Morphogenetic Signaling;

ORGANISM Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
 ; eurosids II; Brassicales; Brassicaceae; Arabidopsids.
 REFERENCE 1 (bases 1 to 65)
 AUTHORS Alonso,J.M., Leisse,T.J., Barajas,P., Chen,H., Cheuk,R., Gadrinab
 , C., Jeske,A., Karnes,M., Kim,C.J., Parker,H., Prednis,L., Shinn,P.,
 Zimmerman,J., and Ecker,J.R.
 TITLE A Sequence-Indexed Library of Insertion Mutations in the
 Arabidopsis Genome
 JOURNAL Unpublished
 COMMENT Contact: Joseph R. Ecker
 Salk Institute Genomic Analysis Laboratory (SIGAL)
 The Salk Institute for Biological Studies
 10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
 Tel: 858 453 4100 x1752
 Fax: 858 558 6379
 Email: ecker@salk.edu
 This is single pass sequence recovered from the left border of
 TDNA. This sequence lies within 300 bases of the 5' end of
 At5g15860.
 Class: TDNA tagged.
 Location/Qualifiers
 1. .65
 /organism="Arabidopsis thaliana"
 /mol_type="genomic DNA"
 /strain="Columbia 0"
 /db_xref="taxon:3702"
 /clone="SALK_059169.41.80.x"
 /clone_lib="Arabidopsis thaliana TDNA insertion lines"
 /notes="PCR was performed on Arabidopsis thaliana lines
 each of which contains one or more TDNA insertion
 elements. The resultant fragment for each line was
 directly sequenced to determine the genomic sequence at
 the site of insertion. Details of the protocols used can
 be found at http://signal.salk.edu/tdna_protocols.html"

BASE COUNT 34 a 5 c 21 g 5 t
 ORIGIN

Query Match 100.0%; Score 10; DB 28; Length 65;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 Db 25 CTTCTCTTTT 16

RESULT 64
 CC179177/c
 LOCUS CC179177
 DEFINITION SALK_059169.41.80.x Arabidopsis thaliana TDNA insertion lines
 Arabidopsis thaliana genomic clone SALK_059169.41.80.x, genomic
 survey sequence.
 ACCESSION CC179177
 VERSION CC179177.1 GI:30317728
 KEYWORDS GSS.
 SOURCE Arabidopsis thaliana (thale cress)

Chuo-ku, Minatojima-minamimachi 2-2-3, Kobe, Hyogo 650-0047, Japan
(E-mail: shayashi@cdb.riken.go.jp, Tel: 81-78-301-3184,
Fax: 81-78-301-3183)

COMMENT This clone was isolated from genomic DNA flanking an insertion of the P element vector P[GaWB] of a *Drosophila* strain.

FEATURES

Location/Qualifiers

1..65

/organism="Drosophila melanogaster"

/mol_type="genomic DNA"

/strain="NP1611"

/db_xref="taxon:7227"

/chromosome="3"

/map="62E8"

/clone="NP1611-5-1"

/note="flanking P[GaWB] transposon insertion"

6 a 34 c 4 g 20 t 1 others

BASE COUNT

ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 65;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

|||||

Db 41 CTTCTCTTTT 50

RESULT 66

AL769814/c

LOCUS

AL769814 65 bp DNA linear GSS 19-JUN-2002

Arabidopsis thaliana T-DNA flanking sequence GK-092H08-012001,

genomic survey sequence.

ACCESSION

AL769814

VERSION

AL769814.1

KEYWORDS

GSS.

SOURCE

Arabidopsis thaliana (thale cress)

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;

rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE

1

Strizhov,N., Li,Y., Rosso,M., Viehoveer,P., Dekker,K., Saedler,H.

and Weisshaar,B.

A pipeline for automated high-throughput generation of ESTs

(flanking sequence tags) from Arabidopsis thaliana T-DNA

transformed lines

Unpublished

JOURNAL

REFERENCE

2

Rosso,M., Strizhov,N., Li,Y., Reiss,B., Dekker,K. and Weisshaar,B.

A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat)

for flanking sequence tag based reverse genetics

Unpublished

JOURNAL

REFERENCE

3 (bases 1 to 65)

Rosso,M., Strizhov,N., Li,Y. and Weisshaar,B.

Submitted (17-JUN-2002) Weisshaar B., Max-Planck-Institut fuer

Zuechtungsforchung, Carl-von-Linne-Weg 10, Koeln 50829, Germany

This sequence is recovered from the right border of the T-DNA. It

indicates an insertion within the locus defined by clone MG46. The

sequences are generated at the MPI for Plant Breeding Research in

the context of the GABI-Kat project. GABI-Kat is part of the German

Plant Genomics program designated 'GABI'. Information on line

availability can be found at:

http://www.mpiz-koeln.mpg.de/GABI-Kat/.

FEATURES

source

1..65

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Columbia 0"

/db_xref="taxon:3702"

/clone="GK-092H08-012001"

/note="PCR was performed on DNA from Arabidopsis thaliana

plants (T1) which were transformed with the T-DNA from vector pAC161. The lines contain one or more T-DNA insertions. The DNA fragment(s) resulting from the PCR were directly sequenced to determine the genomic sequence flanking the insertion. Sequences displaying significant similarity to the A. thaliana nuclear genome sequence were processed for submission. T-DNA derived sequences were removed"

BASE COUNT 32 a 9 c 11 g 13 t

ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 65;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

|||||

Db 39 CTTCTCTTTT 30

RESULT 67

AL949773/c

LOCUS

AL949773 65 bp DNA linear GSS 24-OCT-2002

Arabidopsis thaliana T-DNA flanking sequence GK-323D01-015951,

genomic survey sequence.

ACCESSION

AL949773

VERSION

AL949773.1

KEYWORDS

GSS.

SOURCE

Arabidopsis thaliana (thale cress)

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;

rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE

1

Strizhov,N., Li,Y., Rosso,M., Viehoveer,P., Dekker,K., Saedler,H.

and Weisshaar,B.

A pipeline for automated high-throughput generation of ESTs

(flanking sequence tags) from Arabidopsis thaliana T-DNA

transformed lines

Unpublished

JOURNAL

REFERENCE

2

Rosso,M., Strizhov,N., Li,Y., Reiss,B., Dekker,K. and Weisshaar,B.

A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat)

for flanking sequence tag based reverse genetics

Unpublished

JOURNAL

REFERENCE

3 (bases 1 to 65)

Strizhov,N., Li,Y., Rosso,M. and Weisshaar,B.

Submitted (21-OCT-2002) Weisshaar B., Max-Planck-Institut fuer

Zuechtungsforchung, Carl-von-Linne-Weg 10, Koeln 50829, Germany

This sequence is recovered from the left border of the T-DNA. It

indicates an insertion within the locus defined by clone F9H3. The

sequences are generated at the MPI for Plant Breeding Research in

the context of the GABI-Kat project. GABI-Kat is part of the German

Plant Genomics program designated 'GABI'. Information on line

availability can be found at:

http://www.mpiz-koeln.mpg.de/GABI-Kat/.

FEATURES

source

1..65

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Columbia 0"

/db_xref="taxon:3702"

/clone="GK-323D01-015951"

/note="PCR was performed on DNA from Arabidopsis thaliana

plants (T1) which were transformed with the T-DNA from

vector pAC161. The lines contain one or more T-DNA

insertions. The DNA fragment(s) resulting from the PCR

were directly sequenced to determine the genomic sequence

flanking the insertion. Sequences displaying significant

similarity to the A. thaliana nuclear genome sequence were

processed for submission. T-DNA derived sequences were

```

BASE COUNT      removed"
ORIGIN           26 a    7 c    17 g    15 t

Query Match      100.0%; Score 10; DB 29; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 12 CTTCTCTTTT 3

RESULT 68
TAL29A12P/c
LOCUS TAL29A12P 65 bp DNA linear GSS 13-DEC-2000
DEFINITION T. brucei sheared genomic DNA clone 129a12, forward sequence,
            genomic survey sequence.
ACCESSION AL463978
VERSION AL463978.1 GI:11834241
KEYWORDS GSS.
SOURCE Trypanosoma brucei
ORGANISM Trypanosoma brucei
            Eukaryota; Euklenozoa; Kinetoplastida; Trypanosomatidae;
            Trypanosoma.
REFERENCE 1 (bases 1 to 65)
AUTHORS Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R.,
            Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L.,
            Melville, S.E., Rajandream, M.A. and Barrell, B.G.
TITLE Direct Submission
JOURNAL Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
            project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
            Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and
            nh@sanger.ac.uk
COMMENT Constructed at the Institute for Genomic Research (TIGR),
            Rockville, MD. Genomic DNA isolated from a cloned population of
            Trypanosoma brucei (TREU27/4 GUTat 10.1) was mechanically sheared
            to give a tight size distribution (
            4 kb). The v + i method used for the library construction is
            described in detail in Smith, H. and Venter, J.C. (Making small
            insert libraries for whole genome shotgun sequencing projects. In
            Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.
            Barrell, Oxford University Press, 1999).
            Email: nh@sanger.ac.uk
            Details of T. brucei sequencing at the Sanger Centre are available
            at http://www.sanger.ac.uk/Projects/T_brucei/.
            Location/Qualifiers
                1. .66
                /organism="Trypanosoma brucei"
                /mol_type="genomic DNA"
                /strain="TREU27"
                /db_xref="taxon:5691"
                /clone="129a12"

BASE COUNT      31 a    7 c    22 g    5 t
ORIGIN

Query Match      100.0%; Score 10; DB 29; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 62 CTTCTCTTTT 53

RESULT 69
W85242/c
LOCUS W85242 66 bp mRNA linear EST 12-SEP-1996
DEFINITION m52h08.r1 Soares mouse embryo NBME13.5 14.5 Mus musculus cDNA
            clone IMAGE:408735 5' similar to gb:U13546 rnal Human HMG-17 gene
            for non-histone chromosomal protein (HUMAN); gb:U12944 Mouse mRNA
            for HMG-17 chromosomal protein (MOUSE); mRNA sequence.
            W85242

ACCESSION
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

```

```

VERSION W85242.1 GI:1397731
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 66)
AUTHORS Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,
            Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,
            Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B.,
            Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and
            Waterston, R.
TITLE The WashU-HMI Mouse EST Project
JOURNAL Unpublished
COMMENT Contact: Marra M/Mouse EST Project
            WashU-HMI Mouse EST Project
            Washington University School of Medicine
            4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
            Tel: 314 286 1800
            Fax: 314 286 1810
            Email: mouseest@watson.wustl.edu
            This clone is available royalty-free through LNL; contact the
            IMAGE Consortium (info@image.llnl.gov) for further information.
            MGI:252503
            Trace considered overall poor quality
            Seq primer: -28M13 rev2 from Amerisham
            High quality sequence stop: 1.
            Location/Qualifiers
                1. .66
                /organism="Mus musculus"
                /mol_type="mRNA"
                /strain="C57BL/6J"
                /db_xref="taxon:10090"
                /clone="IMAGE:408735"
                /sex="unknown"
                /tissue_type="embryo"
                /dev_stage="13.5-14.5dpc total fetus"
                /lab_host="DH10B"
                /clone_lib="Soares mouse embryo NBME13.5 14.5"
                /note="vector: pT73D-Pac (Pharmacia) with a modified
                polylinker; Site 1: Not 1; Site 2: Eco RI; 1st strand cDNA
                was primed with a Not I - oligo(gt) primer (5'
                TGTTACCAATCTGAAGTGGGCGGCGGCGGAAATTTTTTTTTTTTTTTT
                T 3'), on equal amounts of mRNA from 2 13.5dpc and 2
                14.5dpc embryos [total RNA provided by Minoru Ko, Wayne
                State Univ., from 2 ]; double-stranded cDNA was ligated to
                Eco RI adaptors (Pharmacia), digested with Not I and
                cloned into the Not I and Eco RI sites of the modified
                pT73 vector. Library went through one round of
                normalization, and was constructed by Bento Soares and
                M.Patima Bonaldo. "
                BASE COUNT      19 a    19 c    18 g    10 t
                ORIGIN

Query Match      100.0%; Score 10; DB 14; Length 66;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
    |||||
Db 41 CTTCTCTTTT 32

RESULT 70
A2514401
LOCUS A2514401
DEFINITION 1M0361X04F Mouse 10kb plasmid UUGCLM library Mus musculus genomic
            clone UUGCLM0361X04 F, genomic survey sequence.
            A2514401
ACCESSION A2514401
VERSION A2514401.1 GI:10695717
KEYWORDS GSS
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

```

Rukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 (bases 1 to 66)

REFERENCE
AUTHORS
 Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C., Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T., Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von Niedehausen, A., and Wright, D., Weiss, R.

TITLE
 Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts

JOURNAL
COMMENT
 Contact: Robert B. Weiss
 University of Utah
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0361 row: K column: 04
 Seq primer: CGTGTAAACGACGGCCAGT
 Class: plasmid ends
 High quality sequence stop: 66.

FEATURES
 Location/Qualifiers
 1..66
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUC1M0361K04"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold. T1-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUC1M library"
 /note="Vector: PWD42nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource (http://www.jax.org/resources/documents/dnares/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of PWD42 [gil4732114|gb|AF129072.1], a copy-number inducible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."

BASE COUNT
 20 a 10 c 8 g 28 t

ORIGIN
 Query Match 100.0%; Score 10; DB 28; Length 66;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
 Db 17 CTTCTCTTTT 26

RESULT 71
 A1442885
 LOCUS
 DEFINITION
 sa28b08.x1 Gm-cl004 Glycine max cDNA clone GENOME SYSTEMS CLONE ID: Gm-cl004-592 3' similar to TR:024482 024482 SAL13-2.; mRNA sequence.

ACCESSION
 A1442885
 VERSION
 A1442885.1 GI:4299305
 KEYWORDS
 EST.

Glycine max (soybean)
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids I; Fabales; Fabaceae; Papilionoideae; Phaseoleae; Glycine. 1 (bases 1 to 67)

REFERENCE
AUTHORS
 Shoemaker, R., Keim, P., Vodkin, L., Erpelding, J., Coryell, V., Khanna, A., Bolla, B., Marra, M., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T., Underwood, K., Steptoe, M., Theising, B., Allen, M., Bowers, Y., Person, B., Swaller, T., Gibbons, M., Pape, D., Harvey, N., Schuck, R., Ritter, E., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R., and Wilson, R.

TITLE
 Public Soybean EST Project

JOURNAL
COMMENT
 Unpublished
 Contact: Shoemaker R/Public Soybean EST Project
 Public Soybean EST Project
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu

Trace considered overall poor quality
 Possible reversed clone: similarity on wrong strand This clone is available through: ResGen, Invitrogen Corp. 2130 South Memorial Parkway Huntsville, AL 35801 For further information call: (800)-533-4363 or contact via email: ccu@resgen.com
 Seq primer: -40UP from Gibco
 High quality sequence stop: 1
 POLYA=No.

FEATURES
 Location/Qualifiers
 1..67
 /organism="Glycine max"
 /mol_type="mRNA"
 /db_xref="taxon:3847"
 /clone="GENOME SYSTEMS CLONE ID: Gm-cl004-592"
 /tissue_type="root"
 /lab_host="XL10-Gold"
 /clone_lib="Gm-cl004"
 /note="Vector: pBluescript II XR; Site 1: EcoRI; Site 2: XhoI; Root cDNA. The mRNA was isolated from entire roots of 8 day old 'Williams' seedlings which were propagated on paper towels with distilled water. Stratagene's cDNA Synthesis Kit (catalog #200401) was used to synthesize the cDNA. First-strand synthesis was performed with 5-methyl dCTP, hence the ligated cDNA is hemimethylated. Stratagene's first-strand synthesis primer was used [CAGAGAGAGAGAGAGAGACTAGTCGAG(T)-18]. After second-strand synthesis, the cDNA ends were 'polished' with clone Pfu DNA polymerase, ligated to EcoRI adaptors, and phosphorylated. The XhoI site within the first-strand synthesis primer was restricted by digestion with XhoI; all XhoI sites in the cDNA would be protected by their hemimethylated status. The cDNA constructs were size-fractionated with a 500bp cutoff, using GibcoBRL Life Technologies' cDNA Size Fractionation column. The column eluent was then ligated into Stratagene's pBluescript II XR Predigested vector (pBluescript II SK(+)) that had been digested with EcoRI and XhoI, and phosphorylated. Both the white and blue colonies appear to contain recombinant plasmids with cDNA inserts. Blue colonies 9n=15) have been sequenced, and possess putative cDNA inserts. This library was constructed by Dr. Paul Keim & Virginia H. Coryell, Department of Biology, Box 5640, Northern Arizona University, Flagstaff, AZ 86011, Phone: 520-523-1078 (Dr. Paul Keim), 520-523-1372 (Virginia H. Coryell), Fax: 520-523-7500, email: paul.keim@naui.edu, virginia.coryell@naui.edu"

BASE COUNT
 18 a 18 c 6 g 24 t 1 others

ORIGIN
 Query Match 100.0%; Score 10; DB 9; Length 67;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||
 Db 56 CTTCTCTTTT 65

RESULT 72
 CD029241
 LOCUS
 DEFINITION
 mgns007xa10f.b Magnaporthe grisea NS Uni-Zap XR Library Magnaporthe
 grisea cDNA clone mgns007xa10 5', mRNA sequence.

ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM

CD029241.1 GI:30410697
 Magnaporthe grisea (anamorph: Pyricularia grisea)
 Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;
 Sordariomycetes incertae sedis; Magnaporthaceae; Magnaporthe.
 1 (bases 1 to 67)
 Ebbola, D.J., Yuan, J., Thomas, T.L., Bobrowicz, P., Lu, G., Bhatterai
 , K. and Dean, R.A.
 Expressed sequence tags from the rice blast fungus, Magnaporthe
 grisea
 Unpublished
 Contact: Ebbola DJ
 Department of Plant Pathology & Microbiology
 Texas A&M University
 Peterson Bldg, MS2132, College Station, TX 77843-2132, USA
 Tel: 979 845 4831
 Fax: 979 845 6483
 Email: d-ebbola@tamu.edu
 Chromatogram file of this sequence is available, see contact person

PCR Primers
 FORWARD: T3 primer
 BACKWARD: T7 primer
 Plate: mgns007 row: A column: 10
 Seq primer: T3.

FEATURES
 source
 1..67
 Location/Qualifiers
 /organism="Magnaporthe grisea"
 /mol_type="mRNA"
 /strain="Guy11"
 /db_xref="taxon:148305"
 /clone="mgns007xa10"
 /sex="Mat1-2 hermaphrodite"
 /cell_type="mycelium"
 /clone_lib="Magnaporthe grisea NS Uni-Zap XR Library"
 /note="Vector: pBluescriptSK-; Site 1: EcoRI; Site 2: XhoI
 ; Unidirectional cloning. EcoRI site has T3 primer and
 predominantly 5' reads. T7 primer on XhoI side of insert.
 Nitrogen starvation library. Cells were inoculated into
 minimal medium and grown for two days with shaking (150
 rpm) at room temperature. Culture was harvested, blended,
 inoculated into minimal medium as above for 24 h. Cells
 were harvested, washed with water and inoculated into
 minimal medium base lacking nitrogen source for 6 h.
 Sequences were processed by one of two methods. Where a
 full-length alignment to the M. grisea genome sequence was
 available, the EST sequence was trimmed according to the
 alignment, otherwise sequence quality was assessed using
 phredPhrap version 991019 and trimmed according to phd
 files (0.05) and for vector seqs."

BASE COUNT
 ORIGIN
 15 a 18 c 14 g 20 t

Query Match 100.0%; Score 10; DB 14; Length 67;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||

Db 23 CTTCTCTTTT 32

RESULT 73
 BM447291/c
 LOCUS
 DEFINITION
 DSA008D10.57203 An expressed sequence tag database for the
 halotolerant green alga, Dunaliella salina Dunaliella salina cDNA
 clone DSA008D10 5, mRNA sequence.

ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM

BM447291.1 GI:19852863
 Dunaliella salina
 Dunaliella salina
 Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
 Dunaliellaceae; Dunaliella.
 1 (bases 1 to 68)
 Cushman, J.C.
 An expressed sequence tag database for the halotolerant green alga,
 Dunaliella salina
 Unpublished
 Contact: Cushman JC
 Department of Biochemistry
 University of Nevada
 MS200, Reno, NV 89557-0014, USA
 Tel: 775-784-1918
 Fax: 775-784-1650
 Email: jcushman@unr.edu
 PCR Primers
 FORWARD: T3 20mer
 BACKWARD: T7 21mer
 Plate: 008 row: D column: 10
 Seq primer: T3 20mer
 High quality sequence stop: 68.

FEATURES
 source
 1..68
 Location/Qualifiers
 /organism="Dunaliella salina"
 /mol_type="mRNA"
 /db_xref="taxon:3046"
 /clone="DSA008D10"
 /tissue_type="Cells, which was adapted in 2.5M NaCl via a
 incremental series from 1.7 to 2.0 to 2.25 to 2.5 M NaCl,
 were exposed to 3.4 M NaCl for 5 hours"
 /cell_type="Green"
 /clone_lib="An expressed sequence tag database for the
 halotolerant green alga, Dunaliella salina"
 /note="Vector: Lambda Uni-Zap XR, Bluescript SK-; Site 1:
 EcoRI; Site 2: XhoI; Library construction was performed
 according to Stratagene's recommended protocol for the
 Lambda Uni-ZapXR vector and cDNA synthesis kit."

BASE COUNT
 ORIGIN
 22 a 13 c 24 g 9 t

Query Match 100.0%; Score 10; DB 12; Length 68;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 |||||
 Db 42 CTTCTCTTTT 33

RESULT 74
 AL757142/c
 LOCUS
 DEFINITION
 Arabidopsis thaliana T-DNA flanking sequence GK-118D11-012821,
 genomic survey sequence.

ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM

AL757142.1 GI:21495490
 GSS.
 Arabidopsis thaliana (thale cress)
 Arabidopsis thaliana
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;

Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE
AUTHORS Strizhov,N., Li,Y., Rosso,M., Viehoveer,P., Dekker,K., Saedler,H. and Weisshaar,B.

TITLE A pipeline for automated high-throughput generation of FSTs (flanking sequence tags) from Arabidopsis thaliana T-DNA transformed lines

JOURNAL Unpublished

REFERENCE
AUTHORS Rosso,M., Strizhov,N., Li,Y., Reiss,B., Dekker,K. and Weisshaar,B.
TITLE A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat) for flanking sequence tag based reverse genetics

JOURNAL Unpublished

REFERENCE
AUTHORS Strizhov,N., Li,Y., Rosso,M. and Weisshaar,B.

JOURNAL Submitted (17-JUN-2002) Weisshaar B., Max-Planck-Institut fuer Zuechtungsforschung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany
TITLE This sequence is recovered from the right border of the T-DNA. It indicates an insertion within the locus defined by clone F27K19. The sequences are generated at the MPI for Plant Breeding Research in the context of the GABI-Kat project. GABI-Kat is part of the German Plant Genomics program designated 'GABI'. Information on line availability can be found at:
http://www.mpiz-koeln.mpg.de/GABI-Kat/.

FEATURES

Location/Qualifiers

1..68
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="GK-118D11-012821"
/note="PCR was performed on DNA from Arabidopsis thaliana (clone lib="Arabidopsis thaliana T-DNA insertion lines" notes="T1) which were transformed with the T-DNA from vector PAC161. The lines contain one or more T-DNA insertions. The DNA fragment(s) resulting from the PCR were directly sequenced to determine the genomic sequence flanking the insertion. Sequences displaying significant similarity to the A. thaliana nuclear genome sequence were processed for submission. T-DNA derived sequences were removed"

BASE COUNT 35 a 8 c 7 g 17 t 1 others

Query Match 100.0%; Score 10; DB 29; Length 68;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

Db 53 CTTCTCTTTT 44

RESULT 75

LOCUS AL651716
DEFINITION AL651716 XGC-gastrula Silurana tropicalis cDNA clone TGas036d16 5', mRNA sequence.
ACCESSION AL651716
VERSION AL651716.1 GI:17661928

KEYWORDS EST.

SOURCE Silurana tropicalis (western clawed frog)

ORGANISM Silurana tropicalis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Amphibia; Batrachia; Anura; Mesobatrachia; Pipidae; Xenopodinae; Silurana.

REFERENCE 1 (bases 1 to 70)

AUTHORS Huckle,E., Taylor,R., Ashurst,J.L., Zorn,A.M. and Rogers,J.

TITLE Sanger Xenopus tropicalis EST project 2001 (10_2001)

JOURNAL Unpublished

COMMENT Contact: Huckle E

Sanger Centre
Hinxton, Cambridgeshire, CB10 1SA, UK
Email: trop@sanger.ac.uk

Sanger Xenopus tropicalis EST project 2001

TROPICALIS_SEQUENCE ID: TGas036d16.sp6

Sequencing primer: SP6

This sequence is from a Xenopus Gene Collection (XGC) library constructed by Aaron M. Zorn.

FEATURES

Location/Qualifiers

1..70
/organism="Silurana tropicalis"
/mol_type="mRNA"
/db_xref="taxon:8364"
/clone="TGas036d16"
/dev_stage="gastrula (stages 10.5-13 mixed)"
/lab_host="Escherichia coli XL1-blue"
/clone_lib="XGC-gastrula"
/note="Vector: pCS107; Site 1: EcoRI; Site 2: NotI; cDNA was oligo dT primed from Sug of poly A+ RNA from stages 10-13 gastrulae. EcoRI-NotI cut cDNA was then ligated into pCS107 with EcoRI at the 5' end and NotI at the 3' end."

BASE COUNT 19 a 17 c 15 g 19 t

ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 70;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10

Db 5 CTTCTCTTTT 14

RESULT 76

LOCUS AA255635
DEFINITION AA255635 70 bp mRNA linear EST 13-AUG-1997
2831507.r1 NCI CGAP GCBI Homo sapiens cDNA clone IMAGE:686821 5' similar to SW:YKUD_YEAST P36042 HYPOPHYSICAL 21.2 KD PROTEIN IN TOR2-PAS1 INTERGENIC REGION.; contains element TARI repetitive element ;, mRNA sequence.
ACCESSION AA255635
VERSION AA255635.1 GI:1892570

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 70)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index

JOURNAL Unpublished

COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaps-remail.nih.gov

This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.

Trace considered overall poor quality

Possible reversed clone: similarity on wrong strand

Insert Length: 1914 Std Error: 0.00

Seq primer: -28m13 rev2 ET from Amersham

High quality sequence stop: 1.

Location/Qualifiers

1..70

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:686821"

/tissue_type="germinal center B cell"

/lab_host="DH10B"

/notes="Vector: pT773D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA

was prepared from human tonsillar cells enriched for germinal center B cells by flow sorting (CD20+, IgD-), provided by Dr. Louis M. Staudt (NCI), Dr. David Allman (NCI) and Dr. Gerald Marti (CBER). cDNA synthesis was primed with a Not I - oligo(dT) primer
 (5'-TGTTACCAATCTGAAGTGGAGCGCGCTCAATTTTTTTTTTTT-3',
 1. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified p773 vector. Library went through one round of normalization, and was constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 0 a 32 c 0 g 38 t
 ORIGIN
 Query Match 100.0%; Score 10; DB 9; Length 70;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
 Db 15 CTTCTCTTTT 24

RESULT 77
 H45651/c
 LOCUS
 DEFINITION Yn97d02.s1 Soares adult brain N2B5HB55Y Homo sapiens cDNA clone IMAGE:176355 3' similar to gb|K01562|HUMCRH1 Human Ro RNA (rRNA); ; mRNA sequence.

ACCESSION H45651.1 GI:921703
 VERSION EST.
 KEYWORDS Homo sapiens (human)
 SOURCE Homo sapiens

ORGANISM
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 70)
 Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevas, E., Waterston, R., Williamson, A., Wohlmann, P. and Wilson, R.
 The WashU-Merck EST Project

TITLE Unpublished
 JOURNAL
 COMMENT Contact: Wilson RK
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu

Insert Size: 979
 High quality sequence starts: 1
 High quality sequence stops: 1
 Source: IMAGE Consortium, LLNL
 This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
 Trace considered overall poor quality
 Insert Length: 979 Std Error: 0.00
 Seq primer: SP6
 High quality sequence stop: 1.

FEATURES
 source
 1..70
 Location/Qualifiers
 /organism="Homo sapiens"
 /mol_type="rRNA"
 /db_xref="GDB:3838551"
 /db_xref="taxon:9606"
 /clone="IMAGE:176355"
 /sex="Male"
 /dev_stage="55-year old"
 /lab_host="DH10B (ampicillin resistant)"
 /clone_lib="Soares adult brain N2B5HB55Y"
 /note="Organ: brain; Vector: p773D (Pharmacia) with a modified polylinker; Site: 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5'

TGTTACCAATCTGAAGTGGAGCGCGCTCAATTTTTTTTTTTT 3',
 double-stranded cDNA was size selected, ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of a modified p773 vector (Pharmacia). Library went through one round of normalization to a Cot = 53. Library constructed by Bento Soares and M. Fatima Bonaldo. The adult brain RNA was provided by Dr. Donald H. Gilden. Tissue was acquired 17-18 hours after death which occurred in consequence of a ruptured aortic aneurysm. RNA was prepared from a pool of tissues representing the following areas of the brain: frontal, parietal, temporal and occipital cortex from the left and right hemispheres, subcortical white matter, basal ganglia, thalamus, cerebellum, midbrain, pons and medulla."

BASE COUNT 27 a 11 c 18 g 14 t
 ORIGIN

Query Match 100.0%; Score 10; DB 14; Length 70;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 |||||
 Db 13 CTTCTCTTTT 4

RESULT 78
 AZ918371/c

LOCUS
 DEFINITION 1006004B08.x1 1006 - RescueMu Grid G Zea mays genomic, genomic survey sequence.

ACCESSION AZ918371
 VERSION AZ918371.1 GI:13387655
 KEYWORDS GSS.
 SOURCE Zea mays

ORGANISM
 Zea mays

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACCAD clade; Panicoideae; Andropogoneae; Zea.
 1 (bases 1 to 70)

REFERENCE 1
 AUTHORS Walbot V.
 TITLE Maize genomic sequences found using engineered RescueMu transposon
 JOURNAL Unpublished
 COMMENT Contact: Walbot V
 Department of Biological Sciences
 Stanford University
 855 California Ave, Palo Alto, CA 94304, USA

Tel: 650 723 2227
 Fax: 650 725 8221
 Email: walbot@stanford.edu
 Plate: 1006004 row: 36
 Class: transposon-tagged.
 Location/Qualifiers
 1..70
 /organism="Zea mays"
 /mol_type="genomic DNA"
 /cultivar="mixed background W23/A188/B73"
 /db_xref="taxon:4577"
 /tissue_type="leaf"
 /dev_stage="adult"
 /lab_host="DH10B"
 /clone_lib="1006 - RescueMu Grid G"
 /note="Organ: leaf; Vector: RescueMu (engineered from Bluescript backbone); Site: 1: BamHI; Site 2: BglII; RescueMu is a 4.9 kb, modified maize Mu transposon designed to allow plasmid rescue from total genomic DNA. Mu elements insert preferentially into transcription units. For more information on RescueMu, go to the web site 'www.zmdb.iastate.edu' and follow the links for 'RescueMu.' Grid G was grown at Stanford in 2000. DNA was extracted from leaf punches, double digested using BamHI and BglII, and ligated to form circular plasmids. DH10B

FEATURES
 source

cells were transformed and then screened on LB plates with ampicillin."

BASE COUNT 23 a 11 c 19 g 17 t

Query Match 100.0%; Score 10; DB 28; Length 70;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 44 CTTCTCTTTT 35

RESULT 79
D78209/c
LOCUS D78209 EST from 8p21.3-p22 Homo sapiens cDNA clone B6-1-5, mRNA
DEFINITION D78209
ACCESSION D78209.1 GI:2104127
VERSION
KEYWORDS
SOURCE
ORGANISM Homo sapiens (human)

REFERENCE
AUTHORS
TITLE
Isolation of 45 exon-like fragments from 8p22-->p21.3, a region that is commonly deleted in hepatocellular, colorectal, and non-small cell lung carcinomas
JOURNAL
MEDLINE
PUBMED
COMMENT
Contact: Yusuke Nakamura
Institute of Medical Science
University of Tokyo
4-6-1, Shirokanedai, Minato-ku, Tokyo 108, Japan
Tel: 81-3-5449-5372
Fax: 81-3-5449-5433
Email: yusuke@ims.u-tokyo.ac.jp.

FEATURES
source
1..71
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/map="8p21.3-p22"
/clone="B6-1-5"
/clone_lib="EST from 8p21.3-p22"
BASE COUNT 29 a 8 c 21 g 13 t

Query Match 100.0%; Score 10; DB 14; Length 71;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 27 CTTCTCTTTT 18

RESULT 80
BZ770025/c
LOCUS BZ770025.1 GI:28943709
DEFINITION
Arabidopsis thaliana genomic clone SALK_142959.51.10.x, genomic survey sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM Arabidopsis thaliana (thale cress)

Query Match 100.0%; Score 10; DB 14; Length 71;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 27 CTTCTCTTTT 18

RESULT 80
BZ770025/c
LOCUS BZ770025.1 GI:28943709
DEFINITION
Arabidopsis thaliana genomic clone SALK_142959.51.10.x, genomic survey sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM Arabidopsis thaliana (thale cress)

Query Match 100.0%; Score 10; DB 29; Length 71;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 69 CTTCTCTTTT 60

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.

REFERENCE
AUTHORS
TITLE
A Sequence-Indexed Library of Insertion Mutations in the Arabidopsis Genome
JOURNAL
COMMENT
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGNAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@sgalk.edu

FEATURES
source
1..71
Location/Qualifiers
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="SALK_142959.51.10.x"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines each of which contains one or more TDNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at http://signal.salk.edu/tdna_protocols.html"
BASE COUNT 28 a 13 c 8 g 22 t

Query Match 100.0%; Score 10; DB 29; Length 71;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 69 CTTCTCTTTT 60

RESULT 81
AA184862/c
LOCUS AA184862.1 GI:1768508
DEFINITION
musculus (house mouse)
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM Mus musculus (house mouse)
REFERENCE
AUTHORS
TITLE
The WashU-HMMI Mouse EST Project
JOURNAL
COMMENT
Contact: Marra M/Mouse EST Project
WashU-HMMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810

Query Match 100.0%; Score 10; DB 29; Length 71;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 69 CTTCTCTTTT 60

RESULT 81
AA184862/c
LOCUS AA184862.1 GI:1768508
DEFINITION
musculus (house mouse)
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM Mus musculus (house mouse)
REFERENCE
AUTHORS
TITLE
The WashU-HMMI Mouse EST Project
JOURNAL
COMMENT
Contact: Marra M/Mouse EST Project
WashU-HMMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810

Query Match 100.0%; Score 10; DB 29; Length 71;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 69 CTTCTCTTTT 60

RESULT 81
AA184862/c
LOCUS AA184862.1 GI:1768508
DEFINITION
musculus (house mouse)
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM Mus musculus (house mouse)
REFERENCE
AUTHORS
TITLE
The WashU-HMMI Mouse EST Project
JOURNAL
COMMENT
Contact: Marra M/Mouse EST Project
WashU-HMMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810

Query Match 100.0%; Score 10; DB 29; Length 71;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 69 CTTCTCTTTT 60

Email: mouseestwatson.wustl.edu
This clone is available royalty-free through LNL ; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.
MGI:394901

Trace considered overall poor quality
Seq primer: -28M13 rev2 from Amersham
High quality sequence stop: 1.

FEATURES

source
Location/Qualifiers
1..72
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="IMAGE:642909"
/sex="male"
/tissue_type="lymph node"
/dev_stage="4 weeks"
/lab_host="DH10B"

/clone_lib="Soares mouse lymph node NbMLN"
/note="organ: lymph node; Vector: pT73D-Pac (Pharmacia)
with a modified polylinker; Site 1: Not 1; Site 2: Eco RI;
1st strand cDNA was primed with a Not I - oligo(dT) primer
[5',
TGTACCAATCTGAAGTGGAGCGCGCGATCTTTTTTTTTTTTTTTTTTTTT
3']; double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pT73 vector. RNA
provided by Dr. Bertrand Jordan. Library constructed and
normalized by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 28 a 13 c 17 g 14 t

ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 72;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
DB 22 CTTCTCTTTT 13

RESULT 82

EX285944 72 bp DNA linear GSS 07-MAR-2003
LOCUS Arabidopsis thaliana T-DNA flanking sequence GK-386H01-018249,
Genomic survey sequence.

ACCESSION EX285944

VERSION EX285944.1 GI:28884940

KEYWORDS GSS

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE 1 Strizhov, N., Li, Y., Rosso, M., Viehoever, P., Dekker, K., Saedler, H.
and Weisshaar, B.

A pipeline for automated high-throughput generation of FSTs
(flanking sequence tags) from Arabidopsis thaliana T-DNA
transformed lines
Unpublished

JOURNAL

REFERENCE

AUTHORS Rosso, M., Strizhov, N., Li, Y., Reiss, B., Dekker, K. and Weisshaar, B.
TITLE A new Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat)
for flanking sequence tag based reverse genetics

JOURNAL Unpublished

REFERENCE 3 (bases 1 to 72)

AUTHORS Strizhov, N., Rosso, M., Li, Y. and Weisshaar, B.

TITLE Direct Submission

JOURNAL Submitted (07-MAR-2003) Weisshaar B., Max-Planck-Institut fuer
Zuechtungsforchung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany
This sequence is recovered from the left border of the T-DNA. It
indicates an insertion close to or within gene At2g28150. The

sequences are generated at the MPI for Plant Breeding Research in
the context of the GABI-Kat project. GABI-Kat is part of the German
Plant Genomics program designated 'GABI'. Information on line
availability can be found at:
http://www.mpiz-koeln.mpg.de/GABI-Kat/.

FEATURES

source
Location/Qualifiers
1..72
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="GK-386H01-018249"
/note="pPCR was performed on DNA from Arabidopsis thaliana
plants (T1) which were transformed with the T-DNA from
vector pAC161. The lines contain one or more T-DNA from
insertions. The DNA fragment(s) resulting from the PCR
were directly sequenced to determine the genomic sequence
flanking the insertion. Sequences displaying significant
similarity to the A. thaliana nuclear genome sequence were
processed for submission. T-DNA derived sequences were
removed"

BASE COUNT 11 a 11 c 16 g 34 t

ORIGIN

Query Match 100.0%; Score 10; DB 29; Length 72;

Best Local Similarity 100.0%; Pred. No. 1.6e+05;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

|||||

DB 17 CTTCTCTTTT 26

RESULT 83

AA262253/C

LOCUS

DEFINITION

AA262253

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

1 (bases 1 to 73)

Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J., Moore, B.,
Schellenberg, K., Steptoe, M., Tan, F., Theising, B., White, Y., Wylie,
T., Waterston, R. and Wilson, R.

WashU-Merck EST Project 1997

Unpublished

COMMENT

Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: est@watson.wustl.edu

This clone is available royalty-free through LNL ; contact the
IMAGE Consortium (info@image.lnl.gov) for further information.

Insert Length: 1520 Std Error: 0.00

Seq primer: -28M13 rev2 ET from Amersham

High quality sequence stop: 65.

Location/Qualifiers

1..73

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:668790"

/tissue_type="Pooled human melanocyte, fetal heart, and
pregnant uterus"

BASE COUNT 24 a 13 c 15 g 21 t

ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 73;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05; Indels 0; Gaps 0;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 Db 33 CTTCTCTTTT 24

RESULT 84
 AW497651/c
 LOCUS
 DEFINITION
 SWYD25CAU11A07SK Brugia malayi young adult day 25 cdNA
 (SAW99MLW-BmyD25) Brugia malayi cDNA clone SWYD25CAU11A07 5', mRNA
 sequence.

ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM

Brugia malayi
 Brugia malayi
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
 Onchocercidae; Brugia.
 1 (bases 1 to 73)
 Williams, S.A.
 Genes expressed in young adult day 25 of Brugia malayi
 Contact: Steven A. Williams
 Smith College Department of Biological Sciences
 Department of Biological Sciences, Clark Science Center, Smith
 College, Northampton, MA, 01063, USA
 Tel: 4135853826
 Fax: 4135853786
 Email: genome@smith.edu
 Seq primer: pBluescript SK.

FEATURES
 source
 1..73
 /organism="Brugia malayi"
 /mol_type="mRNA"
 /db_xref="taxon:6279"
 /clone="SWYD25CAU11A07"
 /dev_stage="young adult, twenty five days after infection"
 /lab_host="XLI-Blue MRF,"
 /clone_lib="Brugia malayi young adult day 25 cdNA
 (SAW99MLW-BmyD25)"
 /note="Vector: Lambda Uni-Zap XR; Site 1: Eco RI; Site 2:
 Xho I; Lymphatic filarial nematode parasite of humans
 mRNA was prepared from young adult worms isolated from
 the peritoneal cavity of jirds on day 25 after infection
 and converted to double-stranded cDNA using reverse
 transcriptase and oligo(dT) followed by RNase H and DNA
 pol I. The library has 6.2 x 105 independent recombinants
 and the average insert size is approx. 1101bp. The library
 was constructed by Michelle Lizotte-Waniewski. The
 library is available from Dr. S.A. Williams, email:
 genome@neal.smith.edu."

BASE COUNT 28 a 12 c 21 g 12 t

ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 73;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05; Indels 0; Gaps 0;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 Db 43 CTTCTCTTTT 34

RESULT 85
 AW600116/c
 LOCUS
 DEFINITION
 SWL4CAK10B02SK Brugia malayi L4 cDNA (SAW99MLW-Bml4) Brugia malayi
 cDNA clone SWL4CAK10B02 5', mRNA sequence.

ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM

Brugia malayi
 Brugia malayi
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
 Onchocercidae; Brugia.
 1 (bases 1 to 73)
 Williams, S.A.
 Genes expressed in fourth stage larvae of Brugia malayi
 Contact: Steven A. Williams
 Smith College Department of Biological Sciences
 Department of Biological Sciences, Clark Science Center, Smith
 College, Northampton, MA, 01063, USA
 Tel: 4135853826
 Fax: 4135853786
 Email: genome@smith.edu
 Seq primer: pBluescript SK.

FEATURES
 source
 1..73
 /organism="Brugia malayi"
 /mol_type="mRNA"
 /db_xref="taxon:6279"
 /clone="SWL4CAK10B02"
 /dev_stage="larval stage four"
 /lab_host="XLI-Blue MRF,"
 /clone_lib="Brugia malayi L4 cDNA (SAW99MLW-Bml4)"
 /note="Vector: Lambda Uni-Zap XR; Site 1: Eco RI; Site 2:
 Xho I; Lymphatic filarial nematode parasite of humans
 mRNA was prepared from L4s isolated from the peritoneal
 cavity of jirds and converted to double-stranded cDNA
 using reverse transcriptase and oligo(dT) followed by
 RNase H and DNA pol I. The library has 2.7 x 105
 independent recombinants and the average insert size is
 approx. 1050bp. The library was constructed by Michelle
 Lizotte-Waniewski. The library is available from Dr. S.A.
 Williams, email: genome@neal.smith.edu."

BASE COUNT 30 a 11 c 20 g 12 t

BASE COUNT 30 a 11 c 20 g 12 t

ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 73;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05; Indels 0; Gaps 0;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
 Db 38 CTTCTCTTTT 29

RESULT 86
 AW626514/c
 LOCUS
 DEFINITION
 SNOWJ3CAN64C12SK Onchocerca volvulus infective larva cDNA
 (SAW94WL-OvJ3) Onchocerca volvulus cDNA clone SNOWJ3CAN64C12 5',
 mRNA sequence.

BASE COUNT 28 a 12 c 21 g 12 t

ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
 Onchocerca volvulus
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
 Onchocercidae; Onchocerca.
 1 (bases 1 to 73)
 Williams, S.A., Lu, W., Lizotte-Waniewski, M. and Laney, S.J.
 Genes expressed in infective third stage larvae of Onchocerca
 volvulus
 Unpublished
 Contact: Steven A. Williams
 Molecular Parasitology
 Smith College Department of Biological Sciences
 Department of Biological Sciences, Clark Science Center, Smith
 College, Northampton, MA, 01063, USA
 Tel: 4135853826
 Fax: 4135853786
 Email: genome@smith.edu
 Seq primer: pBluescript SK.
 Location/Qualifiers
 FEATURES
 source
 1..73
 /organism="Onchocerca volvulus"
 /mol_type="mRNA"
 /strain="Sierra Leone"
 /db_xref="taxon:6282"
 /clones="SMOVL3CAN64C12"
 /lab_host="XLI-Blue MRF"
 /clone_lib="Onchocerca volvulus infective larva cDNA
 (SAM94WL-OvL3)"
 /note="Vector: lambda UniZap XR; Site_1: EcoR I; Site_2:
 Xho I; Cutaneous filarial nematode parasite of humans.
 mRNA was prepared from third stage infective larvae of
 Onchocerca volvulus isolated from mosquitoes 10 days after
 infection and converted to double stranded cDNA using
 reverse transcriptase and oligo(dT) followed by RNase H
 and DNase I. The library had 1.8 x 10E5 independent
 recombinants and average insert size was 900 base pairs.
 The library was constructed by Wenhong Lu. The library is
 available from Dr. S.A. Williams, email genome@smith.edu."
 BASE COUNT 29 a 9 c 22 g 13 t
 ORIGIN
 Query Match 100.0%; Score 10; DB 9; Length 73;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 |||||
 Db 20 CTTCTCTTTT 11
 RESULT 87
 AW626555/c 73 bp mRNA linear EST 30-MAR-2000
 LOCUS
 DEFINITION
 (SAM94WL-OvL3) Onchocerca volvulus cDNA clone SMOVL3CAN66D01 5',
 mRNA sequence.
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
 Onchocerca volvulus
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
 Onchocercidae; Onchocerca.
 1 (bases 1 to 73)
 Williams, S.A., Lu, W., Lizotte-Waniewski, M. and Laney, S.J.
 Genes expressed in infective third stage larvae of Onchocerca
 volvulus
 Unpublished
 Contact: Steven A. Williams
 Molecular Parasitology

Smith College Department of Biological Sciences
 Department of Biological Sciences, Clark Science Center, Smith
 College, Northampton, MA, 01063, USA
 Tel: 4135853826
 Fax: 4135853786
 Email: genome@smith.edu
 Seq primer: pBluescript SK.
 Location/Qualifiers
 FEATURES
 source
 1..73
 /organism="Onchocerca volvulus"
 /mol_type="mRNA"
 /strain="Sierra Leone"
 /db_xref="taxon:6282"
 /clones="SMOVL3CAN66D01"
 /lab_host="XLI-Blue MRF"
 /clone_lib="Onchocerca volvulus infective larva cDNA
 (SAM94WL-OvL3)"
 /note="Vector: lambda UniZap XR; Site_1: EcoR I; Site_2:
 Xho I; Cutaneous filarial nematode parasite of humans.
 mRNA was prepared from third stage infective larvae of
 Onchocerca volvulus isolated from mosquitoes 10 days after
 infection and converted to double stranded cDNA using
 reverse transcriptase and oligo(dT) followed by RNase H
 and DNase I. The library had 1.8 x 10E5 independent
 recombinants and average insert size was 900 base pairs.
 The library was constructed by Wenhong Lu. The library is
 available from Dr. S.A. Williams, email genome@smith.edu."
 BASE COUNT 26 a 11 c 21 g 15 t
 ORIGIN
 Query Match 100.0%; Score 10; DB 9; Length 73;
 Best Local Similarity 100.0%; Pred. No. 1.6e+05;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CTTCTCTTTT 10
 |||||
 Db 15 CTTCTCTTTT 6
 RESULT 89
 AW651817/c 73 bp mRNA linear EST 04-APR-2000
 LOCUS
 DEFINITION
 (SWD25CAU14C02SK Brugia malayi young adult day 25 cDNA
 (SAM99MLW-BmyD25) Brugia malayi cDNA clone SWD25CAU14C02 5', mRNA
 sequence.
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
 Brugia malayi
 Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
 Onchocercidae; Brugia.
 1 (bases 1 to 73)
 Williams, S.A.
 Genes expressed in young adult day 25 of Brugia malayi
 Unpublished
 Contact: Steven A. Williams
 Molecular Parasitology
 Smith College Department of Biological Sciences
 Department of Biological Sciences, Clark Science Center, Smith
 College, Northampton, MA, 01063, USA
 Tel: 4135853826
 Fax: 4135853786
 Email: genome@smith.edu
 Seq primer: pBluescript SK.
 Location/Qualifiers
 FEATURES
 source
 1..73
 /organism="Brugia malayi"
 /mol_type="mRNA"
 /db_xref="taxon:6279"
 /clone="SWD25CAU14C02"
 /dev_stage="young adult, twenty five days after infection"
 /lab_host="XLI-Blue MRF"

```

/clone_lib="Brugia malayi young adult day 25 cDNA
(SAW99MLW-BmYD25)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Lymphatic filarial nematode parasite of humans.
mRNA was prepared from young adult worms isolated from
the peritoneal cavity of jirds on day 25 after infection
and converted to double-stranded cDNA using reverse
transcriptase and oligo(dT) followed by RNase H and DNA
pol I. The library has 6.2 x 105 independent recombinants
and the average insert size is approx.1101bp. The library
was constructed by Michelle Lizotte-Waniewski. The
library is available from Dr. S.A. Williams, email:
genome@neal.smith.edu."
BASE COUNT      29 a 11 c 22 g 11 t
ORIGIN
      100.0%; Score 10; DB 9; Length 73;
      Best Local Similarity 100.0%; Pred. No. 1.6e+05;
      Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
      |||||
Db      38 CTTCTCTTTT 29

RESULT 89
BG310475      73 bp mRNA linear EST 23-FEB-2001
LOCUS      SWOV3MCAM55G11SK Onchocerca volvulus molting L3 larva cDNA
DEFINITION      (SL96MLW-Ovml3) Onchocerca volvulus cDNA clone SWOV3MCAM55G11 5',
mRNA sequence.
ACCESSION      BG310475
VERSION      BG310475.1 GI:13112333
KEYWORDS      EST.
SOURCE      Onchocerca volvulus
ORGANISM      Onchocerca volvulus
REFERENCE      Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
AUTHORS      Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.
TITLE      Genes expressed in molting L3 larvae of Onchocerca volvulus
JOURNAL      Unpublished
COMMENT      Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
FEATURES
    source
    1..73
    /organism="Onchocerca volvulus"
    /mol_type="mRNA"
    /strain="Kumba, Cameroons"
    /db_xref="taxon:6282"
    /clone="SWOV3MCAM55G11"
    /dev_stage="molting L3"
    /lab_host="XLI-Blue MRF"
    /clone_lib="Onchocerca volvulus molting L3 larva cDNA
(SL96MLW-Ovml3)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. Third-stage
larvae, L3, were isolated from infected black flies in
Cameroon (forest strain). The L3 were cultured in 20% FCS
in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in
culture. L3 of O. volvulus molt to fourth-stage larvae by
day 5 in culture. mRNA was isolated from approximately
6000 molting larvae (mL3), 2000 larvae from day 1, 2 or 3
in culture, and converted to double-stranded cDNA using
reverse transcriptase and oligo(dT) followed by RNase H
and DNA pol I. The library was constructed in the lambda

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Uni-Zap XR vector and has 1 x 10E6 independent
recombinants and the average insert size is ~1200 bp. The
library was constructed by Sara Lustigman and Michelle
Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams.
The library is available from Dr. Sara Lustigman (email:
slustigm@nyc.org)."
BASE COUNT      29 a 10 c 19 g 15 t
ORIGIN
      100.0%; Score 10; DB 10; Length 73;
      Best Local Similarity 100.0%; Pred. No. 1.6e+05;
      Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
      |||||
Db      35 CTTCTCTTTT 26

RESULT 90
AW874933/c      73 bp mRNA linear EST 22-MAY-2000
LOCUS      SWYACAL04D03SK Brugia malayi young adult cDNA (SAW99MLW-BmYA)
DEFINITION      Brugia malayi cDNA clone SWYACAL04D03 5', mRNA sequence.
ACCESSION      AW874933
VERSION      AW874933.1 GI:8012644
KEYWORDS      EST.
SOURCE      Brugia malayi
ORGANISM      Brugia malayi
REFERENCE      Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
AUTHORS      Williams, S.A.
TITLE      Genes expressed in young adult of Brugia malayi
JOURNAL      Unpublished
COMMENT      Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
FEATURES
    source
    1..73
    /organism="Brugia malayi"
    /mol_type="mRNA"
    /db_xref="taxon:6279"
    /clone="SWYACAL04D03"
    /dev_stage="young adult"
    /lab_host="XLI-Blue MRF"
    /clone_lib="Brugia malayi young adult cDNA (SAW99MLW-BmYA)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Lymphatic filarial nematode parasite of humans.
mRNA was prepared from young adult worms isolated from the
peritoneal cavity of jirds and converted to
double-stranded cDNA using reverse transcriptase and
oligo(dT) followed by RNase H and DNA pol I. The library
has 6.5 x 104 independent recombinants and the average
insert size is approx. 800bp. The library was constructed
by Michelle Lizotte-Waniewski. The library is available
from Dr. S.A. Williams, email: genome@neal.smith.edu."
BASE COUNT      25 a 10 c 23 g 15 t
ORIGIN
      100.0%; Score 10; DB 10; Length 73;
      Best Local Similarity 100.0%; Pred. No. 1.6e+05;
      Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
      |||||
Db      13 CTTCTCTTTT 4

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RESULT 91
BE420470/c
LOCUS      BE420470              73 bp    mRNA    linear    EST 24-JUL-2000
DEFINITION SWOVl2CAS09D05SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
            Onchocerca volvulus cDNA clone SWOVl2CAS09D05 5', mRNA sequence.
ACCESSION  BE420470
VERSION     BE420470.1  GI:9418296
KEYWORDS    EST.
SOURCE      Onchocerca volvulus
            Onchocerca volvulus
            Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
            Onchocercidae; Onchocerca.
REFERENCE   1  (bases 1 to 73)
AUTHORS     Williams, S.A.
TITLE       Genes expressed in L2 larvae of Onchocerca volvulus
JOURNAL     Unpublished
COMMENT     Contact: Steven A. Williams
            Smith College Department of Biological Sciences
            Department of Biological Sciences, Clark Science Center, Smith
            College, Northampton, MA, 01063, USA
            Tel: 4135853826
            Fax: 4135853786
            Email: genome@smith.edu
            Seq primer: pBluescript SK.
            Location/Qualifiers
                1..73
                /organism="Onchocerca volvulus"
                /mol_type="mRNA"
                /db_xref="taxon:6282"
                /clone="SWOVl2CAS09D05"
                /dev_stage="L2"
                /lab_host="XLI-Blue MRF"
                /clone_lib="Onchocerca volvulus L2 larvae cDNA
                (SAW98MLW-OvL2)"
                /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
                Xho I; Filarial nematode parasite of humans. mRNA was
                prepared from approximately 9,000 L2s isolated from
                infected mosquitoes from Kumba, Cameroon and converted to
                double-stranded cDNA using reverse transcriptase and
                oligo(dT) followed by RNase H and DNA pol I. The library
                has 7.3 x 10E4 independent recombinants and the average
                insert size is approximately 1kb. The library is
                constructed by Michelle Lizotte-Waniewski. The library is
                available from Dr.S.A.Williams, email: genome@smith.edu."
BASE COUNT  29 a  10 c  19 g  15 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY  1  CTTCTCTTTT 10
Db   |||||
35  CTTCTCTTTT 26

RESULT 92
BE420471/c
LOCUS      BE420471              73 bp    mRNA    linear    EST 24-JUL-2000
DEFINITION SWOVl2CAS09D06SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
            Onchocerca volvulus cDNA clone SWOVl2CAS09D06 5', mRNA sequence.
ACCESSION  BE420471
VERSION     BE420471.1  GI:9418297
KEYWORDS    EST.
SOURCE      Onchocerca volvulus
            Onchocerca volvulus
            Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
            Onchocercidae; Onchocerca.
REFERENCE   1  (bases 1 to 73)
AUTHORS     Williams, S.A.

```

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TITLE       Genes expressed in L2 larvae of Onchocerca volvulus
JOURNAL     Unpublished
COMMENT     Contact: Steven A. Williams
            Molecular Parasitology
            Smith College Department of Biological Sciences
            Department of Biological Sciences, Clark Science Center, Smith
            College, Northampton, MA, 01063, USA
            Tel: 4135853826
            Fax: 4135853786
            Email: genome@smith.edu
            Seq primer: pBluescript SK.
            Location/Qualifiers
                1..73
                /organism="Onchocerca volvulus"
                /mol_type="mRNA"
                /db_xref="taxon:6282"
                /clone="SWOVl2CAS09D06"
                /dev_stage="L2"
                /lab_host="XLI-Blue MRF"
                /clone_lib="Onchocerca volvulus L2 larvae cDNA
                (SAW98MLW-OvL2)"
                /note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
                Xho I; Filarial nematode parasite of humans. mRNA was
                prepared from approximately 9,000 L2s isolated from
                infected mosquitoes from Kumba, Cameroon and converted to
                double-stranded cDNA using reverse transcriptase and
                oligo(dT) followed by RNase H and DNA pol I. The library
                has 7.3 x 10E4 independent recombinants and the average
                insert size is approximately 1kb. The library is
                constructed by Michelle Lizotte-Waniewski. The library is
                available from Dr.S.A.Williams, email: genome@smith.edu."
BASE COUNT  29 a  10 c  21 g  13 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY  1  CTTCTCTTTT 10
Db   |||||
19  CTTCTCTTTT 10

RESULT 93
BE420480/c
LOCUS      BE420480              73 bp    mRNA    linear    EST 24-JUL-2000
DEFINITION SWOVl2CAS09F04SK Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
            Onchocerca volvulus cDNA clone SWOVl2CAS09F04 5', mRNA sequence.
ACCESSION  BE420480
VERSION     BE420480.1  GI:9418306
KEYWORDS    EST.
SOURCE      Onchocerca volvulus
            Onchocerca volvulus
            Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
            Onchocercidae; Onchocerca.
REFERENCE   1  (bases 1 to 73)
AUTHORS     Williams, S.A.
TITLE       Genes expressed in L2 larvae of Onchocerca volvulus
JOURNAL     Unpublished
COMMENT     Contact: Steven A. Williams
            Molecular Parasitology
            Smith College Department of Biological Sciences
            Department of Biological Sciences, Clark Science Center, Smith
            College, Northampton, MA, 01063, USA
            Tel: 4135853826
            Fax: 4135853786
            Email: genome@smith.edu
            Seq primer: pBluescript SK.
            Location/Qualifiers
                1..73
                /organism="Onchocerca volvulus"
                /mol_type="mRNA"
                /db_xref="taxon:6282"

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/clone="SWOVL2CAS09F04"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA
(SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site_1: Eco RI; Site_2:
Xho I; Filarial nematode parasite of humans. mRNA was
prepared from approximately 9,000 L2s isolated from
infected mosquitoes from Kumba, Cameroon and converted to
double-stranded cDNA using reverse transcriptase and
oligo(dT) followed by RNase H and DNA pol I. The library
has 7.3 x 10E4 independent recombinants and the average
insert size is approximately 1kb. The library was
constructed by Michelle Lizotte-Waniewski. The library is
available from Dr.S.A.Williams, email: genome@smith.edu."
BASE COUNT      31 a   12 c   16 g   14 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1  CTTCTCTTTT 10
        |||||
Db      41  CTTCTCTTTT 32

RESULT 94
BE638405/c
LOCUS
DEFINITION
Onchocerca volvulus L2 larvae cDNA (SAW98MLW-OvL2)
ACCESSION
BE638405
VERSION
BE638405.1 GI:9937024
KEYWORDS
SOURCE
Onchocerca volvulus
ORGANISM
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.
REFERENCE
1 (bases 1 to 73)
AUTHORS
Williams, S.A.
TITLE
Genes expressed in L2 larvae of Onchocerca volvulus
JOURNAL
Unpublished
COMMENT
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1..73
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/db_xref="taxon:6282"
/clone="SWOVL2CAS15C01"
/dev_stage="L2"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus L2 larvae cDNA
(SAW98MLW-OvL2)"
/note="Vector: Lambda Uni-ZAP XR; Site_1: Eco RI; Site_2:
Xho I; Filarial nematode parasite of humans. mRNA was
prepared from approximately 9,000 L2s isolated from
infected mosquitoes from Kumba, Cameroon and converted to
double-stranded cDNA using reverse transcriptase and
oligo(dT) followed by RNase H and DNA pol I. The library
has 7.3 x 10E4 independent recombinants and the average
insert size is approximately 1kb. The library was
constructed by Michelle Lizotte-Waniewski. The library is
available from Dr.S.A.Williams, email: genome@smith.edu."
BASE COUNT      29 a   11 c   19 g   13 t   1 others

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ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1  CTTCTCTTTT 10
        |||||
Db      27  CTTCTCTTTT 18

RESULT 95
BF228818/c
LOCUS
DEFINITION
Onchocerca volvulus L2 larvae cDNA (SAW94WL-OvL3)
ACCESSION
BF228818
VERSION
BF228818.1 GI:11141183
KEYWORDS
EST.
SOURCE
Onchocerca volvulus
ORGANISM
Onchocerca volvulus
Eukaryota; Metazoa; Nematoda; Chromadorea; Spirurida; Filarioidea;
Onchocercidae; Onchocerca.
REFERENCE
1 (bases 1 to 73)
AUTHORS
Williams, S.A., Lu, W., Lizotte-Waniewski, M. and Laney, S.J.
TITLE
Genes expressed in infective third stage larvae of Onchocerca
volvulus
JOURNAL
Unpublished
COMMENT
Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1..73
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/strain="Sierra Leone"
/db_xref="taxon:6282"
/clone="SWOVL3CAN76D10"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus infective larva cDNA
(SAW94WL-OvL3)"
/note="Vector: lambda UniZap XR; Site_1: EcoR I; Site_2:
Xho I; Cutaneous filarial nematode parasite of humans.
mRNA was prepared from third stage infective larvae of
Onchocerca volvulus isolated from mosquitoes 10 days after
infection and converted to double stranded cDNA using
reverse transcriptase and oligo(dT) followed by RNase H
and DNAPol I. The library had 1.8 x 10E5 independent
recombinants and average insert size was 900 base pairs.
The library was constructed by Wenhong Lu. The library is
available from Dr. S.A. Williams, email genome@smith.edu."
BASE COUNT      23 a   12 c   22 g   16 t
ORIGIN
Query Match      100.0%; Score 10; DB 10; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1  CTTCTCTTTT 10
        |||||
Db      11  CTTCTCTTTT 2

RESULT 96
CB886667/c
LOCUS

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DEFINITION SMOV3MCAM64D11SK Onchocerca volvulus molting L3 larva cDNA
(SL96MLW-Ovml3) Onchocerca volvulus cDNA clone SMOV3MCAM64D11 5',
mRNA sequence.
ACCESSION CB886667.1 GI:30088462
VERSION CB886667.1
KEYWORDS EST.
SOURCE Onchocerca volvulus
ORGANISM Onchocerca volvulus
REFERENCE 1 (bases 1 to 73)
AUTHORS Williams, S.A., Lizotte-Waniewski, M., Laney, S. and Lustigman, S.
TITLE Genes expressed in molting L3 larvae of Onchocerca volvulus
JOURNAL Unpublished
COMMENT Contact: Steven A. Williams
Molecular Parasitology
Smith College Department of Biological Sciences
Department of Biological Sciences, Clark Science Center, Smith
College, Northampton, MA, 01063, USA
Tel: 4135853826
Fax: 4135853786
Email: genome@smith.edu
Seq primer: pBluescript SK.
Location/Qualifiers
1..73
/organism="Onchocerca volvulus"
/mol_type="mRNA"
/strain="Kumba, Cameroons"
/db_xref="taxon:6282"
/clone="SMOV3MCAM64D11"
/dev_stage="molting L3"
/lab_host="XLI-Blue MRF"
/clone_lib="Onchocerca volvulus molting L3 larva cDNA
(SL96MLW-Ovml3)"
/note="Vector: Lambda Uni-ZAP XR; Site 1: Eco RI; Site 2:
Xho I; Filarial nematode parasite of humans. Third-stage
larvae, L3, were isolated from infected black flies in
Cameroon (forest strain). The L3 were cultured in 20% FCS
in IMDM+ NCTC 135 and collected after day 1, 2, or 3 in
culture. L3 of O. volvulus molt to fourth-stage larvae by
day 5 in culture. mRNA was isolated from approximately
6000 molting larvae (mL3). 2000 larvae from day 1, 2 or 3
in culture, and converted to double-stranded cDNA using
reverse transcriptase and oligo(dT) followed by RNase H
and DNA pol I. The library was constructed in the lambda
Uni-ZAP XR vector and has 1 x 106 independent
recombinants and the average insert size is ~1200 bp. The
library was constructed by Sara Lustigman and Michelle
Lizotte-Waniewski in the Laboratory of Dr. S. A. Williams.
The library is available from Dr. Sara Lustigman (email:
slustig@nycb.org)."
29 a 10 c 20 g 14 t
BASE COUNT
ORIGIN
Query Match 100.0%; Score 10; DB 14; Length 73;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 30 CTTCTCTTTT 21

RESULT 97
BZ660902
LOCUS
DEFINITION BZ660902 74 bp DNA linear GSS 31-JAN-2003
SALK 024363.42.35.x Arabidopsis thaliana TDNA insertion lines
Arabidopsis thaliana genomic clone SALK_024363.42.35.x, genomic
survey sequence.
ACCESSION BZ660902
VERSION BZ660902.1 GI:28174049
KEYWORDS GSS.
SOURCE Arabidopsis thaliana (thale cress)

Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids
; eurosids II; Brassicales; Brassicaceae; Arabidopsids.
1 (bases 1 to 74)
Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R., Gadrinab
, C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prednis, L., Shinn, P.
, Zimmerman, J. and Ecker, J.R.
A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
Unpublished
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of
TDNA. This sequence lies within an annotated exon of At5g56780.
Class: TDNA tagged.
Location/Qualifiers
1..74
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="SALK 024363.42.35.x"
/clone_lib="Arabidopsis thaliana TDNA insertion lines"
/note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more TDNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/tdna\_protocols.html"
15 a 14 c 23 g 22 t
BASE COUNT
ORIGIN
Query Match 100.0%; Score 10; DB 29; Length 74;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 21 CTTCTCTTTT 30

RESULT 98
CB49383/c
LOCUS
DEFINITION CB49383 75 bp mRNA linear EST 16-DEC-2002
kll1g03.y1 Ascaris suum embryo SL1 TOPO v1 Ascaris suum cDNA 5',
mRNA sequence.
ACCESSION CB49383
VERSION CB49383.1 GI:27001294
KEYWORDS EST.
SOURCE Ascaris suum (pig roundworm)
ORGANISM Ascaris suum
Eukaryota; Metazoa; Nematoda; Chromadorea; Ascaridida; Ascaridoidea
; Ascarididae; Ascaris.
1 (bases 1 to 75)
McCarter, J., Clifton, S., Chiapelli, B., Pape, D., Martin, J., Wylie, T.
, Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y.
, Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Tsagaris, Shvili, R.
, Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Steptoe
, M., Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S.
, Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and
Wilson, R.
The Washington Univ. Nematode EST Project, 1999
Unpublished
Contact: McCarter JP
The Washington Univ. Nematode EST Project, 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA

```

Tel: 314 286 1800
Fax: 314 286 1810

Email: estewatson.wustl.edu
The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. Oligo(dT)-SL1 PCR based library. Embryo cDNA PCR products of size >400 nucleotides containing SL1 on the 5' end and oligo(dT) on the 3' end were non-directionally cloned into pCRII-TOPO(Invitrogen) following the Topo TA cloning protocol. 30-60 cell embryo material was provided by Dr. Richard Davis of City University of New York Graduate Center, College of Staten Island, Staten Island, NY (redavis@postbox.csi.cuny.edu).
Putative full length read
The vector to vector length is 82
Seq primer: SL1 primer.

FEATURES

Location/Qualifiers

1..75
/organism="Ascaris suum"
/mol_type="mRNA"
/db_xref="taxon:6253"
/dev_stage="30-60 cell embryo"
/lab_host="DH10B"
/clone_lib="Ascaris suum embryo SL1 TOPO v1"
/note="Vector: pCRII-TOPO (Invitrogen); Site 1: EcoRI; Site 2: EcoRI; The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. Oligo(dT)-SL1 PCR based library. Embryo cDNA PCR products of size >400 nucleotides containing SL1 on the 5' end and oligo(dT) on the 3' end were non-directionally cloned into pCRII-TOPO(Invitrogen) following the Topo TA cloning protocol. 30-60 cell embryo material was provided by Dr. Richard Davis of City University of New York Graduate Center, College of Staten Island, Staten Island, NY (redavis@postbox.csi.cuny.edu)."

BASE COUNT 34 a 10 c 14 g 17 t
ORIGIN

Query Match 100.0%; Score 10; DB 14; Length 75;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 58 CTTCTCTTTT 49

RESULT 99
TA77C02P 75 bp DNA linear GSS 13-DEC-2000
LOCUS T. brucei sheared genomic DNA clone 77c02, forward sequence,
genomic survey sequence.
AL460777
AL460777.1 GI:11860526
GSS.
KEYWORDS Trypanosoma brucei
SOURCE Trypanosoma brucei
ORGANISM Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;
Trypanosoma.

REFERENCE 1 (bases 1 to 75)
AUTHORS Hall, N., Bowman, S., Lennard, N.J., Doggett, J., Atkin, R., Chillingworth, C., Ormond, D., Harris, B., El-Sayed, N., Hou, L., Melville, S.B., Rajandream, M.A. and Barrell, B.G.
DIRECT SUBMISSION
Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton, Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and nh@sanger.ac.uk

COMMENT Constructed at the Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of Trypanosoma brucei (TREU27/4 GUTat 10.1) was mechanically sheared to give a tight size distribution (4 kb). The v + i method used for the library construction is described in detail in Smith, H. and Venter, J.C. (Making small

insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaudin and B. Barrell, Oxford University Press, 1999).
Email: nelsayed@tigr.org
Details of T. brucei sequencing at the Sanger Centre are available at http://www.sanger.ac.uk/projects/T_brucei/.

FEATURES

source

1..75
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Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
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Db 19 CTTCTCTTTT 28

RESULT 100
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LOCUS v061d12.r1 Soares mammary gland NBMWG Mus musculus cDNA clone
DEFINITION IMAGE:1054391 5' similar to SW:TRFE_RAT P12346 SEROTRANSFERRIN ; mRNA sequence.
AA615345
AA615345.1 GI:2502573
EST.
KEYWORDS Mus musculus (house mouse)
SOURCE Mus musculus

ORGANISM Eukaryota; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 76)
AUTHORS Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T., Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M., Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B., Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and Waterston, R.
TITLE The WashU-HMMI Mouse EST Project
JOURNAL Unpublished
COMMENT Contact: Marra M/Mouse EST Project
WashU-HMMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@watson.wustl.edu
This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
MG1:585967

Trace considered overall poor quality
Possible reversed clone: similarity on wrong strand
Seq primer: -28m13 rev2 ET from Amersham
High quality sequence stop: 1.
Location/Qualifiers

FEATURES

source

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/note="Organ: mammary gland; Vector: pT73D-Pac (Pharmacia

) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was primed with a Not I - Oligo(dT) primer [5' TGTACCAATCTGAAGTGGAGCGCGCGAATGGTTTTTTTTTTTTTTTTTTT T 3']; double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. RNA provided by Dr. Minoru Ko, Wayne State Univ. Library constructed and normalized by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 22 a 17 c 21 g 16 t

ORIGIN

Query Match 100.0%; Score 10; DB 9; Length 76;
Best Local Similarity 100.0%; Pred. No. 1.6e+05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTCTCTTTT 10
| | | | |
Db 34 CTCTCTTTT 25

Search completed: October 28, 2003, 18:19:28
Job time : 2140 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: October 28, 2003, 15:14:39 ; Search time 80 Seconds
(without alignments)
55.173 Million cell updates/sec

Title: US-09-335-032-71

Perfect score: 10

Sequence: 1 cttctctttt 10

Scoring table: OMIGO_NUC

Gapop 60.0, Gapext 60.0

Searched: 569978 seqs, 220691566 residues

Word size: 0

Total number of hits satisfying chosen parameters: 1139956

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 500 summaries

Database: Issued Patents NA.*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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C 5	10	100.0	21	2	US-08-229-528-16
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C 49	c				Sequence 134, App
C 50	c				Sequence 135, App
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C 163	10	100.0	420	3	US-08-991-890-1	Sequence 1, Appl	c 236	10	100.0	537	4	US-08-874-102-43	Sequence 43, Appl
C 164	10	100.0	421	3	US-09-385-982-49	Sequence 49, Appl	c 237	10	100.0	537	4	US-08-874-102-45	Sequence 45, Appl
C 165	10	100.0	434	4	US-09-280-116-31	Sequence 31, Appl	c 238	10	100.0	537	4	US-08-984-919A-42	Sequence 42, Appl
C 166	10	100.0	438	4	US-09-134-001C-2357	Sequence 2357, Ap	c 239	10	100.0	537	4	US-08-984-919A-43	Sequence 43, Appl
C 167	10	100.0	440	2	US-08-406-057-1	Sequence 1, Appl	c 240	10	100.0	537	4	US-08-984-919A-45	Sequence 45, Appl
C 168	10	100.0	440	3	US-08-958-316-1	Sequence 1, Appl	c 241	10	100.0	542	3	US-08-906-156A-17	Sequence 17, Appl
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ALIGNMENTS

US-08-173-489C-181/c
; Sequence 181, Application US/08173489C
; Patent No. 5861344
; GENERAL INFORMATION:
; APPLICANT: WANG, C. -G.
; APPLICANT: HEPBURN, A. G.
; TITLE OF INVENTION: GENETIC SEQUENCE ASSAY USING DNA
; TITLE OF INVENTION: TRIPLE-STRAND FORMATION.
; NUMBER OF SEQUENCES: 365
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PROFILE DIAGNOSTIC SCIENCES, INC.,
; STREET: 510 EAST 73RD STREET,
; CITY: NEW YORK
; STATE: NEW YORK
; COUNTRY: USA
; ZIP: 10021.
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 inch, 1.44mb storage
COMPUTER: IBM PC/XT/AT
OPERATING SYSTEM: MS-DOS version 6.2
SOFTWARE: Wordperfect Version 5.1
CURRENT APPLICATION DATA: US/08/173,489C
FILING DATE: 22 DEC 1993
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/968,436
FILING DATE: 29 OCT 1992
ATTORNEY/AGENT INFORMATION:
NAME: Handelman, Joseph H.
REGISTRATION NUMBER: 26,179
REFERENCE/DOCKET NUMBER: U9518-6

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INFORMATION FOR SEQ ID NO: 246:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 bases
TYPE: nucleic acid
STRANDEDNESS: single stranded
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: third strand derived from M. luteus
HYPOTHETICAL: yes
ANTI-SENSE: no
PUBLICATION INFORMATION:
RELEVANT RESIDUES IN SEQ ID NO: 246 :FROM 1 TO 12
US-08-173-489C-246

Query Match 100.0%; Score 10; DB 2; Length 12;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 3 CTTCTCTTTT 12

RESULT 3
US-08-628-417-3
Sequence 3, Application US/08628417
Patent No. 5627054
GENERAL INFORMATION:
APPLICANT: GILLESPIE, DAVID
TITLE OF INVENTION: COMPETITOR PRIMER ASYMMETRIC
TITLE OF INVENTION: POLYMERASE CHAIN REACTION
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: U.S. ARMY CHEMICAL AND BIOLOGICAL
ADDRESSEE: DEFENSE COMMAND
STREET: OFFICE OF THE CHIEF COUNSEL (AMSCB-GC)
CITY: ABERDEEN PROVING GROUND
STATE: MARYLAND
COUNTRY: USA
ZIP: 21010-5423
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/628,417
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: BIFFONI, ULYSSES J
REGISTRATION NUMBER: 39,908
REFERENCE/DOCKET NUMBER: DAM 398-94
TELECOMMUNICATION INFORMATION:
TELEPHONE: 410-671-1158
TELEFAX: 410-671-2534
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 20 bases
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: oligodeoxynucleotide
HYPOTHETICAL: NO
ANTI-SENSE: YES
ORIGINAL SOURCE:
ORGANISM: Staphylococcus aureus
US-08-628-417-3

TELECOMMUNICATION INFORMATION:
TELEPHONE: (attorney) (212) 708-1880
TELEFAX: (attorney) (212) 246-8959
INFORMATION FOR SEQ ID NO: 181:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 base pairs
TYPE: nucleic acid
STRANDEDNESS: double stranded
TOPOLOGY: linear
MOLECULE TYPE: genomic DNA
DESCRIPTION: hepatitis B virus adw2 isolate,
nucleotides 807 to 818
HYPOTHETICAL: no
ANTI-SENSE: no
ORIGINAL SOURCE:
ORGANISM: Hepatitis B virus
INDIVIDUAL ISOLATE: adw2
PUBLICATION INFORMATION:
AUTHORS: Valenzuela, P, Quiroga, M, Zaldivar, J,
AUTHORS: Gray, P, Ruter, W J.
TITLE: The nucleotide sequence of
the Hepatitis B viral genome and the
identification of the major viral genes
JOURNAL: In "Animal Virus Genetics", Fields, B N,
Jaenisch, R, Fox C F eds
VOLUME:
PAGES: 57-70
DATE: 1980
RELEVANT RESIDUES IN SEQ ID NO: 181 :FROM 1 TO 12
US-08-173-489C-181

Query Match 100.0%; Score 10; DB 2; Length 12;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 11 CTTCTCTTTT 2

RESULT 2
US-08-173-489C-246
Sequence 246, Application US/08173489C
Patent No. 5861244
GENERAL INFORMATION:
APPLICANT: WANG, C. -G.
APPLICANT: HEPBURN, A. G.
TITLE OF INVENTION: GENETIC SEQUENCE ASSAY USING DNA
TRIPLE-STRAND FORMATION.
NUMBER OF SEQUENCES: 365
CORRESPONDENCE ADDRESS:
ADDRESSEE: PROFILE DIAGNOSTIC SCIENCES, INC.,
STREET: 510 EAST 73RD STREET,
CITY: NEW YORK
STATE: NEW YORK
COUNTRY: USA
ZIP: 10021.
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 inch, 1.44Mb storage
COMPUTER: IBM PC/XT/AT
OPERATING SYSTEM: MS-DOS version 6.2
SOFTWARE: Wordperfect Version 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/173,489C
FILING DATE: 22 DEC 1993
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/969,436
FILING DATE: 29 OCT 1992
ATTORNEY/AGENT INFORMATION:
NAME: Handelman, Joseph H.
REGISTRATION NUMBER: 26,179
REFERENCE/DOCKET NUMBER: U9518-6

```
Query Match      100.0%; Score 10; DB 1; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      8 CTTCTCTTTT 17

RESULT 4
US-09-198-452A-6382/c
; Sequence 6382, Application US/09198452A
; Patent No. 6559294
; GENERAL INFORMATION:
; APPLICANT: Grifpals, R.
; TITLE OF INVENTION: Chlamydia pneumoniae genomic sequence and polypeptides, fragments
; TITLE OF INVENTION: thereof and uses thereof, in particular for the diagnosis, prevention
; TITLE OF INVENTION: and treatment of infection
; FILE REFERENCE: 9710-003-999
; CURRENT APPLICATION NUMBER: US/09/198,452A
; CURRENT FILING DATE: 1998-11-24
; NUMBER OF SEQ ID NOS: 6949
; SEQ ID NO 6382
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Chlamydia pneumoniae
US-09-198-452A-6382

Query Match      100.0%; Score 10; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      17 CTTCTCTTTT 8

RESULT 5
US-08-229-528-16/c
; Sequence 16, Application US/08229528
; Patent No. 5837447
; GENERAL INFORMATION:
; APPLICANT: GORSKI, Jack
; TITLE OF INVENTION: MONITORING AN IMMUNE RESPONSE BY ANALYSIS OF AMPLIFIED IMMUNO
; NUMBER OF SEQUENCES: 51
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Foley & Lardner
; STREET: P. O. Box 1497
; CITY: Madison
; STATE: Wisconsin
; COUNTRY: USA
; ZIP: 53701-1497
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 1.44 Mb storage
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: MS-DOS 3.3
; SOFTWARE: WordPerfect, Version 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/229,528
; FILING DATE: 18-APR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/868,569
; FILING DATE: 15-APR-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Scanlon, William J.
; REGISTRATION NUMBER: 30,136
; REFERENCE/DOCKET NUMBER: 30383/133
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (608) 258-4284
; TELEFAX: (608) 258-4258
; INFORMATION FOR SEQ ID NO: 16:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21 base pairs
```

```
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: Other nucleic acid;
; DESCRIPTION: Synthetic DNA oligonucleotide
US-08-229-528-16

Query Match      100.0%; Score 10; DB 2; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      18 CTTCTCTTTT 9

RESULT 6
US-09-198-484-8/c
; Sequence 8, Application US/09198484
; Patent No. 6162435
; GENERAL INFORMATION:
; APPLICANT: Minion, F. Chris
; APPLICANT: Hsu, Tsungda
; TITLE OF INVENTION: RECOMBINANT MYCOPLASMA HYOPNEUMONIAE VACCINE
; FILE REFERENCE: I9000.028/P028
; CURRENT APPLICATION NUMBER: US/09/198,484
; CURRENT FILING DATE: 1998-11-24
; NUMBER OF SEQ ID NOS: 10
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 8
; LENGTH: 21
; TYPE: DNA
; ORGANISM: primer
US-09-198-484-8

Query Match      100.0%; Score 10; DB 3; Length 21;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      15 CTTCTCTTTT 6

RESULT 7
US-08-173-489C-65
; Sequence 65, Application US/08173489C
; Patent No. 5861244
; GENERAL INFORMATION:
; APPLICANT: WANG, C. -G.
; APPLICANT: HEBURN, A. G.
; TITLE OF INVENTION: GENETIC SEQUENCE ASSAY USING DNA
; TITLE OF INVENTION: TRIPLE-STRAND FORMATION.
; NUMBER OF SEQUENCES: 365
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PROFILE DIAGNOSTIC SCIENCES, INC.,
; STREET: 510 EAST 73RD STREET,
; CITY: NEW YORK
; STATE: NEW YORK
; COUNTRY: USA
; ZIP: 10021.
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5 inch, 1.44Mb storage
; COMPUTER: IBM PC/XT/AT
; OPERATING SYSTEM: MS-DOS version 6.2
; SOFTWARE: Wordperfect Version 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/173,489C
; FILING DATE: 22 DEC 1993
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/968,436
; FILING DATE: 29 OCT 1992
```

```
;
;
; ATTORNEY/AGENT INFORMATION:
; NAME: Handelman, Joseph H.
; REGISTRATION NUMBER: 26,179
; REFERENCE/DOCKET NUMBER: U9518-6
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (attorney) (212) 708-1880
; TELEFAX: (attorney) (212) 246-8959
; INFORMATION FOR SEQ ID NO: 65:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 22 base pairs
; TYPE: Nucleic Acid
; STRANDEDNESS: double stranded
; TOPOLOGY: linear
; MOLECULE TYPE: Genomic DNA
; DESCRIPTION: gamma-crystallin gene exon 3
; DESCRIPTION: (Accession # K03006) nucleotides 9 to 30
; HYPOTHEICAL: No
; ANTI-SENSE: No
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; POSITION IN GENOME:
; CHROMOSOME/SEGMENT: chromosome 2
; MAP POSITION: 2q33-q35
; PUBLICATION INFORMATION:
; AUTHORS: Meakin, S O, Breitman, M L, Tsui, L C.
; TITLE: Structural and evolutionary
; TITLE: relationships among five members of the human
; TITLE: gamma-crystallin gene family
; JOURNAL: Molecular and Cellular Biology
; VOLUME: 5
; PAGES: 1408-1414
; DATE: 1985
; RELEVANT RESIDUES IN SEQ ID NO: 65 :FROM 1 TO 22
; US-08-173-489C-65

Query Match 100.0%; Score 10; DB 2; Length 22;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 6 CTTCTCTTTT 15

RESULT 8
US-08-488-212A-17/c
; Sequence 17, Application US/08488212A
; Patent No. 5665355
; GENERAL INFORMATION:
; APPLICANT: Primi, Daniele
; TITLE OF INVENTION: Diagnosis and Treatment of
; TITLE OF INVENTION: AIDS Onset
; NUMBER OF SEQUENCES: 57
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Thomas E. Popovich, Thomas
; ADDRESSEE: Popovich & Associates
; STREET: 80 South 8th Street
; CITY: Minneapolis
; STATE: Minnesota
; COUNTRY: USA
; ZIP: 55402-2111
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 1.44 MB
; MEDIUM TYPE: Storage
; COMPUTER: IBM Compatible Compaq Prolinea
; COMPUTER: 4/66
; OPERATING SYSTEM: MS-DOS Version 5
; SOFTWARE: Microsoft Word for Windows
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/488,212A
; FILING DATE: 07-Jun-1995
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/973,485

;
;
; ATTORNEY/AGENT INFORMATION:
; FILING DATE: No. 5665355 September 9, 1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Thomas E. Popovich
; REGISTRATION NUMBER: 30099
; REFERENCE/DOCKET NUMBER: 3678
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (612) 334-8991
; TELEFAX: (612) 334-8994
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 24 bases
; TYPE: Nucleic Acid
; STRANDEDNESS: Single
; TOPOLOGY: Linear
; MOLECULE TYPE: Other nucleic acid
; MOLECULE TYPE: (oligonucleotide useful in amplification of T Cell Receptor
; MOLECULE TYPE: Vb region)
; HYPOTHEICAL: No
; ORIGINAL SOURCE: Synthesized using
; PUBLICATION INFORMATION:
; AUTHORS: Imberti, Luisa; Sottini,
; AUTHORS: Alessandra; Bettinardi, Alessandra; Puoti, Massimo; Primi,
; AUTHORS: Daniele
; TITLE: Selective Depletion in HIV Infection
; TITLE: of T Cells That Bear Specific T Cell Receptor Vb Sequences
; JOURNAL: Science
; VOLUME: 254
; ISSUE: 5033
; PAGES: 860-862
; PUBLICATION DATE: No. 5665355 September 8, 1991
; US-08-488-212A-17

Query Match 100.0%; Score 10; DB 1; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

RESULT 9
US-08-320-306-17/c
; Sequence 17, Application US/08320306
; Patent No. 5891623
; GENERAL INFORMATION:
; APPLICANT: Primi, Daniele
; TITLE OF INVENTION: Diagnosis and Treatment of
; TITLE OF INVENTION: AIDS Onset
; NUMBER OF SEQUENCES: 57
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Thomas E. Popovich, Thomas
; ADDRESSEE: Popovich & Associates
; STREET: 80 South 8th Street
; CITY: Minneapolis
; STATE: Minnesota
; COUNTRY: USA
; ZIP: 55402-2111
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 1.44 MB
; MEDIUM TYPE: Storage
; COMPUTER: IBM Compatible Compaq Prolinea
; COMPUTER: 4/66
; OPERATING SYSTEM: MS-DOS Version 5
; SOFTWARE: Microsoft Word for Windows
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/320,306
; FILING DATE: 06-OCT-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/973,485
```

```

; FILING DATE: No. 5891623ember 9, 1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Thomas E. Popovich
; REGISTRATION NUMBER: 30099
; REFERENCE/DOCKET NUMBER: 3678
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (612) 334-8991
; TELEFAX: (612) 334-8994
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 24 bases
; TYPE: Nucleic Acid
; STRANDEDNESS: Single
; TOPOLOGY: Linear
; MOLECULE TYPE: Other nucleic acid
; MOLECULE TYPE: (oligonucleotide useful in amplification of T Cell Receptor
; MOLECULE TYPE: Vb region)
; HYPOTHETICAL: No
; ORIGINAL SOURCE: Synthesized using
; PUBLICATION INFORMATION:
; AUTHORS: Imberti, Luisa; Sottini,
; AUTHORS: Alessandra; Bettinardi, Alessandra; Puoti, Massimo; Primi,
; AUTHORS: Daniele
; TITLE: Selective Depletion in HIV Infection
; TITLE: of T Cells That Bear Specific T Cell Receptor Vb Sequences
; JOURNAL: Science
; VOLUME: 254
; ISSUE: 5033
; PAGES: 860-862
; PUBLICATION DATE: No. 5891623ember 8, 1991
; US-08-320-306-17

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Query Match 100.0%; Score 10; DB 2; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Qy 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

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RESULT 10
US-08-488-209B-17/c
; Sequence 17, Application US/08488209B
; Patent No. 5925513
; GENERAL INFORMATION:
; APPLICANT: Primi, Daniele
; TITLE OF INVENTION: Diagnosis and Treatment of
; TITLE OF INVENTION: AIDS Onset
; NUMBER OF SEQUENCES: 57
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Thomas E. Popovich, Thomas
; ADDRESSEE: Popovich & Associates
; STREET: 80 South 8th Street
; CITY: Minneapolis
; STATE: Minnesota
; COUNTRY: USA
; ZIP: 55402-2111
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 1.44 MB
; MEDIUM TYPE: Storage
; COMPUTER: IBM Compatible Compaq Prolinea
; COMPUTER: 4/66
; OPERATING SYSTEM: MS-DOS Version 5
; SOFTWARE: Microsoft word for Windows
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/488.209B
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/973,485
; FILING DATE: No. 5925513ember 9, 1992

```

```

; ATTORNEY/AGENT INFORMATION:
; NAME: Thomas E. Popovich
; REGISTRATION NUMBER: 30099
; REFERENCE/DOCKET NUMBER: 3678
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (612) 334-8991
; TELEFAX: (612) 334-8994
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 24 bases
; TYPE: Nucleic Acid
; STRANDEDNESS: Single
; TOPOLOGY: Linear
; MOLECULE TYPE: Other nucleic acid
; MOLECULE TYPE: (oligonucleotide useful in amplification of T Cell Receptor
; MOLECULE TYPE: Vb region)
; HYPOTHETICAL: No
; ORIGINAL SOURCE: Synthesized using
; PUBLICATION INFORMATION:
; AUTHORS: Imberti, Luisa; Sottini,
; AUTHORS: Alessandra; Bettinardi, Alessandra; Puoti, Massimo; Primi,
; AUTHORS: Daniele
; TITLE: Selective Depletion in HIV Infection
; TITLE: of T Cells That Bear Specific T Cell Receptor Vb Sequences
; JOURNAL: Science
; VOLUME: 254
; ISSUE: 5033
; PAGES: 860-862
; PUBLICATION DATE: No. 5925513ember 8, 1991
; US-08-488-209B-17

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Query Match 100.0%; Score 10; DB 2; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Qy 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

```

```

RESULT 11
US-08-408-011-17/c
; Sequence 17, Application US/08408011
; Patent No. 5928642
; GENERAL INFORMATION:
; APPLICANT: Primi, Daniele
; TITLE OF INVENTION: Diagnosis and Treatment of
; TITLE OF INVENTION: AIDS Onset
; NUMBER OF SEQUENCES: 57
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Thomas E. Popovich, Thomas
; ADDRESSEE: Popovich & Associates
; STREET: 80 South 8th Street
; CITY: Minneapolis
; STATE: Minnesota
; COUNTRY: USA
; ZIP: 55402-2111
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 1.44 MB
; MEDIUM TYPE: Storage
; COMPUTER: IBM Compatible Compaq Prolinea
; COMPUTER: 4/66
; OPERATING SYSTEM: MS-DOS Version 5
; SOFTWARE: Microsoft word for Windows
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/408.011
; FILING DATE: 18-OCT-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/973,485
; FILING DATE: No. 5928642ember 9, 1992
; ATTORNEY/AGENT INFORMATION:

```


NAME: Thomas E. Popovich
REGISTRATION NUMBER: 30099
REFERENCE/DOCKET NUMBER: 3678
TELEPHONE: (612) 334-8991
TELEFAX: (612) 334-8994
INFORMATION FOR SEQ ID NO: 17:
SEQUENCE CHARACTERISTICS:
LENGTH: 24 bases
TYPE: Nucleic Acid
STRANDEDNESS: Single
TOPOLOGY: Linear
MOLECULE TYPE: Other nucleic acid
MOLECULE TYPE: (oligonucleotide useful in amplification of T Cell Receptor
MOLECULE TYPE: Vb region)
HYPOTHETICAL: No
ORIGINAL SOURCE: Synthesized using
PUBLICATION INFORMATION:
AUTHORS: Imberti, Luisa; Sottini,
AUTHORS: Alessandra; Bettinardi, Alessandra; Puoti, Massimo; Primi,
AUTHORS: Daniele
TITLE: Selective Depletion in HIV Infection
TITLE: of T Cells That Bear Specific T Cell Receptor Vb Sequences
JOURNAL: Science
VOLUME: 254
ISSUE: 5033
PAGES: 860-862
PUBLICATION DATE: No. 5928642ember 8, 1991
US-08-408-011-17

Query Match 100.0%; Score 10; DB 2; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

RESULT 12
US-08-559-205-16/c
Sequence 16, Application US/08559205
Patent No. 6087096
GENERAL INFORMATION:
APPLICANT: Dau, Peter C.
APPLICANT: Liu, Debang
TITLE OF INVENTION: Method of Intrafamily Fragment Analysis of the T
TITLE OF INVENTION: Cell Receptor ' and Chain CDR3 Regions
NUMBER OF SEQUENCES: 61
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray & Borun
STREET: 6300 Sears Tower, 233 South Wacker Drive
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606-6402
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/559,205
FILING DATE:
CLASSIFICATION: 436
ATTORNEY/AGENT INFORMATION:
NAME: Gass, David A.
REGISTRATION NUMBER: 38,153
REFERENCE/DOCKET NUMBER: 28721/32972
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448

TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 24 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-559-205-16

Query Match 100.0%; Score 10; DB 3; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

RESULT 13
US-09-417-722-3/c
Sequence 3, Application US/09417722
Patent No. 6309837
GENERAL INFORMATION:
APPLICANT: Dean, Ralph A.
APPLICANT: Wang, Yi-Hong
TITLE OF INVENTION: PCR-based Method for Identifying a Fusarium
TITLE OF INVENTION: Wilt-Resistant Genotype in Plants
FILE REFERENCE: PCR ID: Fusarium-resistant genotype
CURRENT APPLICATION NUMBER: US/09/417,722
CURRENT FILING DATE: 1999-10-13
NUMBER OF SEQ ID NOS: 4
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 3
LENGTH: 24
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: forward PCR
OTHER INFORMATION: primer of FM primer pair
US-09-417-722-3

Query Match 100.0%; Score 10; DB 4; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
Db 23 CTTCTCTTTT 14

RESULT 14
5336598-16/c
Patent No. 5336598
APPLICANT: KOTZIN, BRIAN L.; MARRACK, PHILIPPA; KAPPLER,
JOHN; CHOI, YOUNGWON
TITLE OF INVENTION: METHOD FOR DIAGNOSING A SUPERANTIGEN
CAUSED PATHOLOGICAL CONDITION VIA ASSAY OF T-CELLS
NUMBER OF SEQUENCES: 25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/437,370
FILING DATE: 15-NOV-1989
SEQ ID NO: 16
LENGTH: 24
5336598-16

Query Match 100.0%; Score 10; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

Query Match 100.0%; Score 10; DB 6; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
| | | | | | | | | |
Db 13 CTTCTCTTTT 4

RESULT 17
US-08-412-376-23/c
; Sequence 23, Application US/08412376
; Patent No. 5849900
; GENERAL INFORMATION:
; APPLICANT: Moelling, Karin
; TITLE OF INVENTION: Inhibition Of Viruses By Antisense
; TITLE OF INVENTION: Oligomers Capable Of Binding To Polypurine-Rich Tract Of Sin
; TITLE OF INVENTION: Stranded RNA Or RNA-DNA Hybrids
; NUMBER OF SEQUENCES: 42
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Woodcock Washburn Kurtz
; ADDRESSEE: Mackiewicz & No. 5849900ris
; STREET: One Liberty Place - 46th Floor
; CITY: Philadelphia
; STATE: PA
; COUNTRY: USA
; ZIP: 19103
; COMPUTER READABLE FORM:
; MEDIUM TYPE: DISKETTE, 3.5 INCH, 1.44 Mb STORAGE
; COMPUTER: IBM PS/2
; OPERATING SYSTEM: PC-DOS
; SOFTWARE: WORDPERFECT 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/412.376
; FILING DATE: Herewith
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/954,184
; FILING DATE: 29-SEP-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Doreen Yanko Trujillo
; REGISTRATION NUMBER: 35,719
; REFERENCE/DOCKET NUMBER: APOL-0021
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (215) 568-3100
; TELEFAX: (215) 568-3439
; INFORMATION FOR SEQ ID NO: 23:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27
; TYPE: Nucleic Acid
; STRANDEDNESS: Single
; TOPOLOGY: Linear
; ANTI-SENSE: Yes

Query Match 100.0%; Score 10; DB 2; Length 27;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
| | | | | | | | | |
Db 22 CTTCTCTTTT 13

RESULT 18
US-08-584-040-7049/c
; Sequence 7049, Application US/08584040
; Patent No. 6346398
; GENERAL INFORMATION:
; APPLICANT: Pavco, Pamela
; APPLICANT: McSwiggen, James
; APPLICANT: Stinchcomb, Dan T.
; APPLICANT: Escobedo, Jaime

RESULT 15
US-08-628-417-4
; Sequence 4, Application US/08628417
; Patent No. 5627054
; GENERAL INFORMATION:
; APPLICANT: GILLESPIE, DAVID
; TITLE OF INVENTION: COMPETITOR PRIMER ASYMMETRIC
; TITLE OF INVENTION: POLYMERASE CHAIN REACTION
; NUMBER OF SEQUENCES: 7
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: U.S. ARMY CHEMICAL AND BIOLOGICAL
; ADDRESSEE: DEFENSE COMMAND
; STREET: OFFICE OF THE CHIEF COUNSEL (AMSCB-GC)
; CITY: ABERDEEN PROVING GROUND
; STATE: MARYLAND
; COUNTRY: USA
; ZIP: 21010-5423
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/628,417
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: BIFFONI, ULYSSES J
; REGISTRATION NUMBER: 39,908
; REFERENCE/DOCKET NUMBER: DAM 398-94
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 410-671-1158
; TELEFAX: 410-671-2534
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 25 bases
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: oligodeoxynucleotide
; HYPOTHETICAL: NO
; ANTI-SENSE: YES

Query Match 100.0%; Score 10; DB 1; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
| | | | | | | | | |
Db 8 CTTCTCTTTT 17

RESULT 16
5217891-9/c
; Patent No. 5217891
; APPLICANT: BRAKE, ANTHONY J.;VAN DEN BERG, JOHAN A.
; TITLE OF INVENTION: DNA CONSTRUCTS CONTAINING A KLUYVEROMYCES
; A FACTOR LEADER SEQUENCE FOR DIRECTING SECRETION OF HETEROLOGOUS
; POLYPEPTIDES
; NUMBER OF SEQUENCES: 23
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/507,398
; FILING DATE: 09-APR-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 78,551
; FILING DATE: 28-JUL-1987
; SEQ ID NO: 9:
; LENGTH: 25
5217891-9

```

; TITLE OF INVENTION: METHOD AND REAGENT FOR THE
; TITLE OF INVENTION: TREATMENT OF DISEASES OR
; TITLE OF INVENTION: CONDITIONS RELATED TO LEVELS
; TITLE OF INVENTION: OF VASCULAR ENDOTHELIAL
; TITLE OF INVENTION: GROWTH FACTOR
; NUMBER OF SEQUENCES: 8502
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Lyon & Lyon
; STREET: 633 West Fifth Street
; CITY: Los Angeles
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 90071-2066
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
; MEDIUM TYPE: storage
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: IBM P.C. DOS 5.0
; SOFTWARE: Word Perfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/584,040
; FILING DATE: January 11, 1996
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/005,974
; FILING DATE: October 26, 1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Warburg, Richard J.
; REGISTRATION NUMBER: 32,327
; REFERENCE/DOCKET NUMBER: 218/064
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (213) 489-1600
; TELEFAX: (213) 955-0440
; TELEX: 67-3510
; INFORMATION FOR SEQ ID NO: 7049:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 27 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; FEATURE:
; OTHER INFORMATION: The letter "N" represents the stem II region
; OTHER INFORMATION: of an HH ribozyme.
US-08-584-040-7049

Query Match 100.0%; Score 10; DB 4; Length 27;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 27 CTTCTCTTTT 18

RESULT 19
US-09-270-542-133
; Sequence 133, Application US/09270542
; Patent No. 6322976
; GENERAL INFORMATION:
; APPLICANT: Aitman, Timothy
; APPLICANT: Stanton, James
; TITLE OF INVENTION: Compositions and Methods of Disease Diagnosis and
; TITLE OF INVENTION: Therapy
; FILE REFERENCE: 4198/78179
; CURRENT APPLICATION NUMBER: US/09/270,542
; CURRENT FILING DATE: 1999-03-17
; EARLIER APPLICATION NUMBER: 09/221,222
; EARLIER FILING DATE: 1999-12-23
; NUMBER OF SEQ ID NOS: 207
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 133

Query Match 100.0%; Score 10; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17

RESULT 20
US-09-270-542-132
; Sequence 132, Application US/09270542
; Patent No. 6322976
; GENERAL INFORMATION:
; APPLICANT: Aitman, Timothy
; APPLICANT: Stanton, James
; TITLE OF INVENTION: Compositions and Methods of Disease Diagnosis and
; TITLE OF INVENTION: Therapy
; FILE REFERENCE: 4198/78179
; CURRENT APPLICATION NUMBER: US/09/270,542
; CURRENT FILING DATE: 1999-03-17
; EARLIER APPLICATION NUMBER: 09/221,222
; EARLIER FILING DATE: 1999-12-23
; NUMBER OF SEQ ID NOS: 207
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 132
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Rattus norvegicus
US-09-270-542-132

Query Match 100.0%; Score 10; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 8 CTTCTCTTTT 17

RESULT 21
US-09-270-542-134
; Sequence 134, Application US/09270542
; Patent No. 6322976
; GENERAL INFORMATION:
; APPLICANT: Aitman, Timothy
; APPLICANT: Stanton, James
; TITLE OF INVENTION: Compositions and Methods of Disease Diagnosis and
; TITLE OF INVENTION: Therapy
; FILE REFERENCE: 4198/78179
; CURRENT APPLICATION NUMBER: US/09/270,542
; CURRENT FILING DATE: 1999-03-17
; EARLIER APPLICATION NUMBER: 09/221,222
; EARLIER FILING DATE: 1999-12-23
; NUMBER OF SEQ ID NOS: 207
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 134
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Rattus norvegicus
US-09-270-542-134

Query Match 100.0%; Score 10; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY      1 CTTCTCTTTT 10
      |||||
Db      8 CTTCTCTTTT 17

RESULT 22
US-09-270-542-135
; Sequence 135, Application US/09270542
; Patent No. 6322976
; GENERAL INFORMATION:
; APPLICANT: Altman, Timothy
; APPLICANT: Scott, James
; APPLICANT: Stanton, Lawrence
; TITLE OF INVENTION: Compositions and Methods of Disease Diagnosis and
; FILE REFERENCE: 4198/78179
; CURRENT APPLICATION NUMBER: US/09/270,542
; CURRENT FILING DATE: 1999-03-17
; EARLIER APPLICATION NUMBER: 09/221,222
; EARLIER FILING DATE: 1999-12-23
; NUMBER OF SEQ ID NOS: 207
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 135
; LENGTH: 30
; TYPE: DNA
; ORGANISM: Rattus norvegicus
US-09-270-542-135

Query Match      100.0%; Score 10; DB 4; Length 30;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
      |||||
Db      8 CTTCTCTTTT 17

RESULT 23
US-08-961-083-359/c
; Sequence 359, Application US/08961083
; Patent No. 6159469
; GENERAL INFORMATION:
; APPLICANT: Choi et. al.
; TITLE OF INVENTION: Streptococcus pneumoniae Antigens and Vaccines
; NUMBER OF SEQUENCES: 452
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: Maryland
; COUNTRY: USA
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage
; COMPUTER: HP Vectra 486/33
; OPERATING SYSTEM: MSDOS version 6.2
; SOFTWARE: ASCII Text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/961,083
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Brooks, A. Anders
; REGISTRATION NUMBER: 36,373
; REFERENCE/DOCKET NUMBER: PB340P2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (301) 309-8504
; TELEFAX: (301) 309-8512
; INFORMATION FOR SEQ ID NO: 359:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 33 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 359:

US-08-961-083-359

Query Match      100.0%; Score 10; DB 4; Length 33;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
      |||||
Db      23 CTTCTCTTTT 14

RESULT 24
US-09-536-784-359/c
; Sequence 359, Application US/09536784
; Patent No. 6573082
; GENERAL INFORMATION:
; APPLICANT: Choi et. al.
; TITLE OF INVENTION: Streptococcus pneumoniae Antigens and Vaccines
; NUMBER OF SEQUENCES: 452
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: Maryland
; COUNTRY: USA
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage
; COMPUTER: HP Vectra 486/33
; OPERATING SYSTEM: MSDOS version 6.2
; SOFTWARE: ASCII Text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/536,784
; FILING DATE: 30-Oct-1997
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/961,083
; FILING DATE: OCT-30-1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Michelle S. Marks
; REGISTRATION NUMBER: 41,971
; REFERENCE/DOCKET NUMBER: PB340P3
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (301) 309-8504
; TELEFAX: (301) 309-8512
; INFORMATION FOR SEQ ID NO: 359:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 33 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 359:

US-09-536-784-359

Query Match      100.0%; Score 10; DB 4; Length 33;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
      |||||
Db      23 CTTCTCTTTT 14

RESULT 25
US-09-371-772B-10906/c
; Sequence 10906, Application US/0937172B
; Patent No. 6566127
; GENERAL INFORMATION:
; APPLICANT: Ribozyne Pharmaceuticals, Inc.
; SEQUENCE CHARACTERISTICS:
; LENGTH: 33 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 359:
```

```
; APPLICANT: Pavco, Pam
; APPLICANT: McSwigen, Jim
; APPLICANT: Stinchcomb, Dan
; APPLICANT: Escobedo, Jaime
; TITLE OF INVENTION: Method and Reagent for the Treatment of Diseases or Conditions Re
; FILE REFERENCE: MBHB00,876-J (237/198)
; CURRENT APPLICATION NUMBER: US/09/371,772B
; PRIOR FILING DATE: 1999-08-10
; PRIOR APPLICATION NUMBER: US 60/005,974
; PRIOR FILING DATE: 1995-10-26
; PRIOR APPLICATION NUMBER: US 08/584,040
; PRIOR FILING DATE: 1996-01-08
; NUMBER OF SEQ ID NOS: 14225
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 10906
; LENGTH: 38
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
US-09-371-772B-10906

Query Match          100.0%; Score 10; DB 4; Length 38;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 38 CTTCTCTTTT 29

RESULT 26
5217891-13/c
; Patent No. 5217891
; APPLICANT: BRAKE, ANTHONY J.; VAN DEN BERG, JOHAN A.
; TITLE OF INVENTION: DNA CONSTRUCTS CONTAINING A KLUYVEROMYCES
; A FACTOR LEADER SEQUENCE FOR DIRECTING SECRETION OF HETEROLOGOUS
; POLYPEPTIDES
; NUMBER OF SEQUENCES: 23
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/507,398
; FILING DATE: 09-APR-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 78,551
; FILING DATE: 28-JUL-1987
; SEQ ID NO:13:
; LENGTH: 39
5217891-13

Query Match          100.0%; Score 10; DB 6; Length 39;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

RESULT 27
US-09-277-016-20
; Sequence 20, Application US/09277016
; Patent No. 6143529
; GENERAL INFORMATION:
; APPLICANT: Lapidus, Stanley N
; APPLICANT: Shuber, Anthony P
; TITLE OF INVENTION: Methods for improving sensitivity and specificity of
; screening assays
; FILE REFERENCE: EXT-030
; CURRENT APPLICATION NUMBER: US/09/277,016
; CURRENT FILING DATE: 1999-03-26
; EARLIER APPLICATION NUMBER: 08/876,857
; EARLIER FILING DATE: 1997-06-16
```

```
; EARLIER APPLICATION NUMBER: 08/700,583
; EARLIER FILING DATE: 1996-08-14
; NUMBER OF SEQ ID NOS: 37
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 20
; LENGTH: 40
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence:PCR-G-FOR (p53)
; OTHER INFORMATION: Exon 8)
US-09-277-016-20

Query Match          100.0%; Score 10; DB 3; Length 40;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 31 CTTCTCTTTT 40

RESULT 28
US-07-908-317-33
; Sequence 33, Application US/07908317
; Patent No. 5420027
; GENERAL INFORMATION:
; APPLICANT: FISHER, CHARLES W.
; APPLICANT: BARNES, HENRY J.
; APPLICANT: ESTABROOK, RONALD W.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; TITLE OF INVENTION: THE EXPRESSION OF BIOLOGICALLY
; TITLE OF INVENTION: ACTIVE FUSION PROTEINS COMPRISING A
; TITLE OF INVENTION: EUKARYOTIC CYTOCHROME P450 FUSED TO
; TITLE OF INVENTION: A REDUCTASE IN BACTERIA
; NUMBER OF SEQUENCES: 34
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: ARNOLD, WHITE & DURKEE
; STREET: P.O. BOX 4433
; CITY: HOUSTON
; STATE: TEXAS
; COUNTRY: USA
; ZIP: 77210
; COMPUTER READABLE FORM:
; MEDIUM TYPE: FLOPPY DISK
; COMPUTER: IBM PC COMPATIBLE
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: WORDPERFECT 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/908,317
; FILING DATE: 19920702
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: PARKER, DAVID L.
; REGISTRATION NUMBER: 32,165
; REFERENCE/DOCKET NUMBER: UTSD:292/PAR
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 512-320-7200
; TELEFAX: 512-474-7577
; TELEX: NOT APPLICABLE
; INFORMATION FOR SEQ ID NO: 33:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 42 base pairs
; TYPE: NUCLEIC ACID
; STRANDEDNESS: single
; TOPOLOGY: linear
US-07-908-317-33

Query Match          100.0%; Score 10; DB 1; Length 42;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
```

```
Db      25 CTTCTCTTTT 34
|||||
RESULT 29
US-09-461-697-227/c
; Sequence 227, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo. Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Paranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; TITLE OF INVENTION: CELL DEATH
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 227
; LENGTH: 42
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-227

Query Match      100.0%; Score 10; DB 3; Length 42;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
|||||
Db      14 CTTCTCTTTT 5

RESULT 30
PCT-US93-06171-33
; Sequence 33, Application PC/TUS9306171
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: FUSION PROTEINS COMPRISING
; TITLE OF INVENTION: EUKARYOTIC CYTOCHROME P450 FUSED TO
; TITLE OF INVENTION: A REDUCTASE
; NUMBER OF SEQUENCES: 34
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: ARNOLD, WHITE & DURKEE
; STREET: P.O. BOX 4433
; CITY: HOUSTON
; STATE: TEXAS
; COUNTRY: USA
; ZIP: 77210
; COMPUTER READABLE FORM:
; MEDIUM TYPE: FLOPPY DISK
; COMPUTER: IBM PC COMPATIBLE
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: WORDPERFECT 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US93/06171
; FILING DATE: 19930629
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/908,317
; FILING DATE: 02 July 1992
; ATTORNEY/AGENT INFORMATION:
; NAME: PARKER, DAVID L.
; REGISTRATION NUMBER: 32,165
; REFERENCE/DOCKET NUMBER: UTSD:292/PAR
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 512-320-7200
```

```
; TELEFAX: 512-474-7577
; TELEX: NOT APPLICABLE
; INFORMATION FOR SEQ ID NO: 33:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 42 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
PCT-US93-06171-33

Query Match      100.0%; Score 10; DB 5; Length 42;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
|||||
Db      25 CTTCTCTTTT 34

RESULT 31
5258302-15/c
; Patent No. 5258302
; APPLICANT: VEDVICK, THOMAS S.; ENGEL, MICHAEL E.; URCAN,
; MARY S.; BUCKHOLZ, RICHARD G.; KINNEY, JENNIFER A.
; TITLE OF INVENTION: DNA FOR EXPRESSION OF APROTININ IN
; METHYLOTROPHIC YEAST CELLS
; NUMBER OF SEQUENCES: 31
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/560,618
; FILING DATE: 30-JUL-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 547,985
; FILING DATE: 03-JUL-1990
; SEQ ID NO:15;
; LENGTH: 43
5258302-15

Query Match      100.0%; Score 10; DB 6; Length 43;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
|||||
Db      19 CTTCTCTTTT 10

RESULT 32
US-09-671-317-765
; Sequence 765, Application US/09671317
; Patent No. 6528260
; GENERAL INFORMATION:
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; APPLICANT: Bougueleret, Lydie
; APPLICANT: Cohen, Annick
; TITLE OF INVENTION: BIALLELIC MARKERS RELATED TO GENES INVOLVED IN DRUG METABOLISM
; FILE REFERENCE: 62.US3.CIP
; CURRENT APPLICATION NUMBER: US/09/671,317
; CURRENT FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US 09/536,178
; PRIOR FILING DATE: 2000-03-23
; PRIOR APPLICATION NUMBER: PCT/IB00/00403
; PRIOR FILING DATE: 2000-03-24
; PRIOR APPLICATION NUMBER: US 60/126,269
; PRIOR FILING DATE: 1999-03-25
; PRIOR APPLICATION NUMBER: US 60/131,961
; PRIOR FILING DATE: 1999-04-30
; NUMBER OF SEQ ID NOS: 977
; SOFTWARE: Patent.pm
; SEQ ID NO 765
; LENGTH: 47
; TYPE: DNA
; ORGANISM: Homo Sapiens
```

```

; LENGTH: 47
; TYPE: DNA
; ORGANISM: Homo Sapiens
; NAME/KEY: allele
; LOCATION: 24
; FEATURE:
; OTHER INFORMATION: 12-602-196 : polymorphic base C or T
US-09-671-317-765

Query Match      100.0%; Score 10; DB 4; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 1 CTTCTCTTTT 10

RESULT 33
US-09-422-978-522/c
; Sequence 522, Application US/09422978
; Patent No. 6537751
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; TITLE OF INVENTION: Biallelic markers for use in constructing a high density...
; FILE REFERENCE: GENSET.020Cp1
; CURRENT APPLICATION NUMBER: US/09/422,978
; CURRENT FILING DATE: 1999-10-20
; EARLIER APPLICATION NUMBER: US 09/298,850
; EARLIER FILING DATE: 1998-04-21
; EARLIER APPLICATION NUMBER: US 60/109,732
; EARLIER FILING DATE: 1998-11-23
; EARLIER APPLICATION NUMBER: US 60/082,614
; EARLIER FILING DATE: 1998-04-21
; NUMBER OF SEQ ID NOS: 11796
; SEQ ID NO 522
; LENGTH: 47
; TYPE: DNA
; ORGANISM: Homo Sapiens
; NAME/KEY: allele
; LOCATION: 24
; OTHER INFORMATION: 99-15101-154 : polymorphic base G or C
US-09-422-978-522

Query Match      100.0%; Score 10; DB 4; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 15 CTTCTCTTTT 6

RESULT 34
US-09-422-978-1282
; Sequence 1282, Application US/09422978
; Patent No. 6537751
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; TITLE OF INVENTION: Biallelic markers for use in constructing a high density...
; FILE REFERENCE: GENSET.020Cp1
; CURRENT APPLICATION NUMBER: US/09/422,978
; CURRENT FILING DATE: 1999-10-20
; EARLIER APPLICATION NUMBER: US 09/298,850
; EARLIER FILING DATE: 1999-04-21
; EARLIER APPLICATION NUMBER: US 60/109,732
; EARLIER FILING DATE: 1998-11-23
; EARLIER APPLICATION NUMBER: US 60/082,614
; EARLIER FILING DATE: 1998-04-21
; NUMBER OF SEQ ID NOS: 11796
; SEQ ID NO 1282
; LENGTH: 47
; TYPE: DNA
; ORGANISM: Homo Sapiens
; NAME/KEY: allele
; LOCATION: 24
; OTHER INFORMATION: 99-4541-39 : polymorphic base G or T
US-09-422-978-1282

Query Match      100.0%; Score 10; DB 4; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 9 CTTCTCTTTT 18

RESULT 36
US-09-422-978-2472/c
; Sequence 2472, Application US/09422978
; Patent No. 6537751
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; TITLE OF INVENTION: Biallelic markers for use in constructing a high density...
; FILE REFERENCE: GENSET.020Cp1
; CURRENT APPLICATION NUMBER: US/09/422,978
; CURRENT FILING DATE: 1999-10-20
; EARLIER APPLICATION NUMBER: US 09/298,850
; EARLIER FILING DATE: 1999-04-21
; EARLIER APPLICATION NUMBER: US 60/109,732
; EARLIER FILING DATE: 1998-11-23
; EARLIER APPLICATION NUMBER: US 60/082,614
; EARLIER FILING DATE: 1998-04-21
; NUMBER OF SEQ ID NOS: 11796
; SEQ ID NO 1546
; LENGTH: 47
; TYPE: DNA
; ORGANISM: Homo Sapiens
; NAME/KEY: allele
; LOCATION: 24
; OTHER INFORMATION: 99-4541-39 : polymorphic base G or T
US-09-422-978-1546

Query Match      100.0%; Score 10; DB 4; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
   |||||
Db 9 CTTCTCTTTT 18

```

Wed Oct 29 15:38:01 2003

```
; EARLIER FILING DATE: 1998-04-21
; NUMBER OF SEQ ID NOS: 11796
; SEQ ID NO 2472
; LENGTH: 47
; TYPE: DNA
; ORGANISM: Homo Sapiens
; FEATURE:
; NAME/KEY: allele
; LOCATION: 24
; OTHER INFORMATION: 99-11191-86 : polymorphic base A or G
US-09-422-978-2472

Query Match      100.0%; Score 10; DB 4; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 45 CTTCTCTTTT 36

RESULT 37
US-09-422-978-3284/c
; Sequence 3284, Application US/09422978
; Patent No. 6537751
; GENERAL INFORMATION:
; APPLICANT: Cohen, Daniel
; APPLICANT: Blumenfeld, Marta
; APPLICANT: Chumakov, Ilya
; TITLE OF INVENTION: Biallelic markers for use in constructing a high density...
; FILE REFERENCE: GENSET.020CP1
; CURRENT APPLICATION NUMBER: US/09/422,978
; CURRENT FILING DATE: 1999-10-20
; EARLIER APPLICATION NUMBER: US 09/298,850
; EARLIER FILING DATE: 1999-04-21
; EARLIER APPLICATION NUMBER: US 60/109,732
; EARLIER FILING DATE: 1998-11-23
; EARLIER APPLICATION NUMBER: US 60/082,614
; EARLIER FILING DATE: 1998-04-21
; NUMBER OF SEQ ID NOS: 11796
; SEQ ID NO 3284
; LENGTH: 47
; TYPE: DNA
; ORGANISM: Homo Sapiens
; FEATURE:
; NAME/KEY: allele
; LOCATION: 24
; OTHER INFORMATION: 99-2981-53 : polymorphic base T or C
US-09-422-978-3284

Query Match      100.0%; Score 10; DB 4; Length 47;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 41 CTTCTCTTTT 32

RESULT 38
US-09-461-697-225/c
; Sequence 225, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
```

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; TITLE OF INVENTION: CELL DEATH
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 225
; LENGTH: 48
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-225

Query Match      100.0%; Score 10; DB 3; Length 48;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 20 CTTCTCTTTT 11

RESULT 39
US-08-477-928A-37/c
; Sequence 37, Application US/08477928A
; Patent No. 6207389
; GENERAL INFORMATION:
; APPLICANT: Dosch, Hans M.
; TITLE OF INVENTION: METHODS FOR CONTROLLING T
; TITLE OF INVENTION: LYMPHOCYTE MEDIATED IMMUNE RESPONSES
; NUMBER OF SEQUENCES: 49
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BAKER & BOTTS
; STREET: 1299 Pennsylvania Avenue
; CITY: Washington D.C.
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 20004-2400
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/477,928A
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 536
; ATTORNEY/AGENT INFORMATION:
; NAME: Remenick, James
; REGISTRATION NUMBER: 36902
; REFERENCE/DOCKET NUMBER: 19060-0105
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 639 7700
; TELEFAX: (202) 639 7890
; INFORMATION FOR SEQ ID NO: 37:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 50 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-477-928A-37

Query Match      100.0%; Score 10; DB 3; Length 50;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 42 CTTCTCTTTT 33

RESULT 40
US-08-910-632-40/c
```



```
; Sequence 40, Application US/08910632B
; Patent No. 6077668
; GENERAL INFORMATION:
; APPLICANT: KOOL, ERIC T.
; TITLE OF INVENTION: HIGHLY SENSITIVE MULTIMERIC NUCLEIC ACID PROBES
; FILE REFERENCE: 220.00010130
; CURRENT APPLICATION NUMBER: US/08/910,632B
; CURRENT FILING DATE: 1997-08-13
; EARLIER APPLICATION NUMBER: 08/805,631
; EARLIER FILING DATE: 1997-02-26
; EARLIER APPLICATION NUMBER: 08/393,439
; EARLIER FILING DATE: 1995-02-23
; EARLIER APPLICATION NUMBER: 08/047,860
; EARLIER FILING DATE: 1993-04-15
; NUMBER OF SEQ ID NOS: 83
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 40
; LENGTH: 53
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: DNA 53mer circle
US-08-910-632-40

Query Match 100.0%; Score 10; DB 3; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16

RESULT 41
US-08-910-632-41
; Sequence 41, Application US/08910632B
; Patent No. 6077668
; GENERAL INFORMATION:
; APPLICANT: KOOL, ERIC T.
; TITLE OF INVENTION: HIGHLY SENSITIVE MULTIMERIC NUCLEIC ACID PROBES
; FILE REFERENCE: 220.00010130
; CURRENT APPLICATION NUMBER: US/08/910,632B
; CURRENT FILING DATE: 1997-08-13
; EARLIER APPLICATION NUMBER: 08/805,631
; EARLIER FILING DATE: 1997-02-26
; EARLIER APPLICATION NUMBER: 08/393,439
; EARLIER FILING DATE: 1995-02-23
; EARLIER APPLICATION NUMBER: 08/047,860
; EARLIER FILING DATE: 1993-04-15
; NUMBER OF SEQ ID NOS: 83
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 41
; LENGTH: 53
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: stem-loop RNA multimer which binds HIV-1 gag RNA
US-08-910-632-41

Query Match 100.0%; Score 10; DB 3; Length 53;
Best Local Similarity 30.0%; Pred. No. 1.9e+03;
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 38 CUUCUCUUU 47

RESULT 42
US-08-805-631A-40/c
; Sequence 40, Application US/08805631A
; Patent No. 6096880
; GENERAL INFORMATION:
; APPLICANT: UNIVERSITY OF ROCHESTER
; TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND
; NUMBER OF SEQUENCES: 72
; CORRESPONDENCE ADDRESS:
; ADDRESS: MUETING, RAASCH & GEBHARDT, P.A.
; STREET: 119 No. 6096880th Fourth Street, Suite 201
; CITY: Minneapolis
; STATE: Minnesota
; COUNTRY: USA
; ZIP: 55401
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
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```
; APPLICANT: UNIVERSITY OF ROCHESTER
; TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND
; NUMBER OF SEQUENCES: 72
; CORRESPONDENCE ADDRESS:
; ADDRESS: MUETING, RAASCH & GEBHARDT, P.A.
; STREET: 119 No. 6096880th Fourth Street, Suite 201
; CITY: Minneapolis
; STATE: Minnesota
; COUNTRY: USA
; ZIP: 55401
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/805,631A
; FILING DATE: 26-FEB-97
; CLASSIFICATION: 536
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/393,439
; FILING DATE: 23-FEB-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/047,860
; FILING DATE: 15-APR-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: SANDBERG, VICTORIA A.
; REGISTRATION NUMBER: 41,287
; REFERENCE/DOCKET NUMBER: 220.00010140
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 612-305-1226
; TELEFAX: 612-305-1228
; INFORMATION FOR SEQ ID NO: 40:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 53 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
US-08-805-631A-40

Query Match 100.0%; Score 10; DB 3; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16

RESULT 43
US-08-805-631A-41
; Sequence 41, Application US/08805631A
; Patent No. 6096880
; GENERAL INFORMATION:
; APPLICANT: UNIVERSITY OF ROCHESTER
; TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND
; NUMBER OF SEQUENCES: 72
; CORRESPONDENCE ADDRESS:
; ADDRESS: MUETING, RAASCH & GEBHARDT, P.A.
; STREET: 119 No. 6096880th Fourth Street, Suite 201
; CITY: Minneapolis
; STATE: Minnesota
; COUNTRY: USA
; ZIP: 55401
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
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APPLICATION NUMBER: US/08/805,631A
FILING DATE: 26-FEB-97
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/393,439
FILING DATE: 23-FEB-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/047,860
FILING DATE: 15-APR-1993
ATTORNEY/AGENT INFORMATION:
NAME: SANDBERG, VICTORIA A.
REGISTRATION NUMBER: 41,287
REFERENCE/DOCKET NUMBER: 220.00010140
TELEPHONE: 612-305-1226
TELEFAX: 612-305-1228
INFORMATION FOR SEQ ID NO: 41:
SEQUENCE CHARACTERISTICS:
LENGTH: 53 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: RNA (genomic)
US-08-805-631A-41

Query Match 100.0%; Score 10; DB 3; Length 53;
Best Local Similarity 30.0%; Pred. No. 1.9e+03;
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|::|:|::|
DB 38 CUUCUCUUU 47

RESULT 44
US-09-569-344-40/c
Sequence 40, Application US/09569344
Patent No. 6368802
GENERAL INFORMATION:
APPLICANT: UNIVERSITY OF ROCHESTER
TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND
DNA
NUMBER OF SEQUENCES: 72
CORRESPONDENCE ADDRESS:
ADDRESS: MUETING, RAASCH & GEBHARDT, P.A.
STREET: 119 No. 6368802th Fourth Street, Suite 201
CITY: Minneapolis
STATE: Minnesota
COUNTRY: USA
ZIP: 55401
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/569,344
FILING DATE: 11-May-2000
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/805,631
FILING DATE: 26-FEB-97
APPLICATION NUMBER: US 08/393,439
FILING DATE: 23-FEB-1995
APPLICATION NUMBER: US 08/047,860
FILING DATE: 15-APR-1993
ATTORNEY/AGENT INFORMATION:
NAME: SANDBERG, VICTORIA A.
REGISTRATION NUMBER: 41,287
REFERENCE/DOCKET NUMBER: 220.00010140
TELECOMMUNICATION INFORMATION:
TELEPHONE: 612-305-1226
TELEFAX: 612-305-1228

INFORMATION FOR SEQ ID NO: 40:
SEQUENCE CHARACTERISTICS:
LENGTH: 53 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: circular
MOLECULE TYPE: DNA (genomic)
SEQUENCE DESCRIPTION: SEQ ID NO: 40:
US-09-569-344-40

Query Match 100.0%; Score 10; DB 4; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|::|:|::|
DB 25 CTTCTCTTTT 16

RESULT 45
US-09-569-344-41
Sequence 41, Application US/09569344
Patent No. 6368802
GENERAL INFORMATION:
APPLICANT: UNIVERSITY OF ROCHESTER
TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND
DNA
NUMBER OF SEQUENCES: 72
CORRESPONDENCE ADDRESS:
ADDRESS: MUETING, RAASCH & GEBHARDT, P.A.
STREET: 119 No. 6368802th Fourth Street, Suite 201
CITY: Minneapolis
STATE: Minnesota
COUNTRY: USA
ZIP: 55401
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/569,344
FILING DATE: 11-May-2000
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/805,631
FILING DATE: 26-FEB-97
APPLICATION NUMBER: US 08/393,439
FILING DATE: 23-FEB-1995
APPLICATION NUMBER: US 08/047,860
FILING DATE: 15-APR-1993
ATTORNEY/AGENT INFORMATION:
NAME: SANDBERG, VICTORIA A.
REGISTRATION NUMBER: 41,287
REFERENCE/DOCKET NUMBER: 220.00010140
TELECOMMUNICATION INFORMATION:
TELEPHONE: 612-305-1226
TELEFAX: 612-305-1228
INFORMATION FOR SEQ ID NO: 41:
SEQUENCE CHARACTERISTICS:
LENGTH: 53 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: RNA (genomic)
SEQUENCE DESCRIPTION: SEQ ID NO: 41:
US-09-569-344-41

Query Match 100.0%; Score 10; DB 4; Length 53;
Best Local Similarity 30.0%; Pred. No. 1.9e+03;
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10

```

Db          38 CUUCUCUUU 47
;
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Lyon & Lyon
; STREET: 633 West Fifth Street
; CITY: Los Angeles
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 90071
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
; MEDIUM TYPE: storage
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: IBM P.C. DOS 5.0
; SOFTWARE: FastSEQ Version 1.5
; CURRENT APPLICATION DATA:
; FILING DATE: January 16, 1996
; APPLICATION NUMBER: US/08/585,684B
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/000,951
; FILING DATE: July 7, 1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Warburg, Richard
; REGISTRATION NUMBER: 32,327
; REFERENCE/DOCKET NUMBER: 218/078
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (213) 489-1600
; TELEFAX: (213) 955-0440
; TELEX: 67-3510
; INFORMATION FOR SEQ ID NO: 2710:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 54 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
;
; US-08-585-684B-2710
;
; Query Match 100.0%; Score 10; DB 2; Length 54;
; Best Local Similarity 100.0%; Pred. No. 1.9e+03;
; Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
;
; QY 1 CTTCTCTTTT 10
;      |||||
; Db 13 CTTCTCTTTT 4
;
; RESULT 48
; US-09-038-073-2533/c
; Sequence 2533, Application US/09038073
; Patent No. 6194150
;
; GENERAL INFORMATION:
; APPLICANT: Stinchcomb, Daniel T.
; APPLICANT: Jarvis, Thale
; APPLICANT: McSwiggen, James
; TITLE OF INVENTION: METHOD AND REAGENT FOR THE
; TITLE OF INVENTION: INDUCTION OF GRAFT TOLERANCE
; TITLE OF INVENTION: AND REVERSAL OF IMMUNE RESPONSES
; NUMBER OF SEQUENCES: 2751
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Lyon & Lyon
; STREET: 633 West Fifth Street
; CITY: Los Angeles
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 90071
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
; MEDIUM TYPE: storage
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: IBM P.C. DOS 5.0
; SOFTWARE: FastSEQ Version 1.5
; CURRENT APPLICATION DATA:
; FILING DATE: January 16, 1996
; APPLICATION NUMBER: 60/000,951
; FILING DATE: July 7, 1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Warburg, Richard
; REGISTRATION NUMBER: 32,327
; REFERENCE/DOCKET NUMBER: 218/078
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (213) 489-1600
; TELEFAX: (213) 955-0440
; TELEX: 67-3510
; INFORMATION FOR SEQ ID NO: 2533:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 54 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
;
; US-08-585-684B-2533
;
; Query Match 100.0%; Score 10; DB 2; Length 54;
; Best Local Similarity 100.0%; Pred. No. 1.9e+03;
; Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
;
; QY 1 CTTCTCTTTT 10
;      |||||
; Db 13 CTTCTCTTTT 4
;
; RESULT 47
; US-08-585-684B-2710/c
; Sequence 2710, Application US/08585684B
; Patent No. 5877021
;
; GENERAL INFORMATION:
; APPLICANT: Stinchcomb, Daniel T.
; APPLICANT: Jarvis, Thale
; APPLICANT: McSwiggen, James
; TITLE OF INVENTION: METHOD AND REAGENT FOR THE
; TITLE OF INVENTION: INDUCTION OF GRAFT TOLERANCE
; TITLE OF INVENTION: AND REVERSAL OF IMMUNE RESPONSES
; NUMBER OF SEQUENCES: 2751

```

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;
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/585,684
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Warburg, Richard
; REGISTRATION NUMBER: 32,327
; REFERENCE/DOCKET NUMBER: 218/078
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (213) 489-1600
; TELEFAX: (213) 955-0440
; TELEX: 67-3510
; INFORMATION FOR SEQ ID NO: 2533:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 54 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-09-038-073-2533

Query Match 100.0%; Score 10; DB 3; Length 54;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

RESULT 49
US-09-038-073-2710/c
; Sequence 2710, Application US/09038073
; Patent No. 6194150
; GENERAL INFORMATION:
; APPLICANT: Stinchcomb, Daniel T.
; APPLICANT: McSwigen, James
; TITLE OF INVENTION: METHOD AND REAGENT FOR THE
; TITLE OF INVENTION: INDUCTION OF GRAFT TOLERANCE
; TITLE OF INVENTION: AND REVERSAL OF IMMUNE RESPONSES
; NUMBER OF SEQUENCES: 2751
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Lyon & Lyon
; STREET: 633 West Fifth Street
; STREET: Suite 4700
; CITY: Los Angeles
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 90071
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
; MEDIUM TYPE: storage
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: IBM P.C. DOS 5.0
; SOFTWARE: FastSeq Version 1.5
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/038,073
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/585,684
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Warburg, Richard
; REGISTRATION NUMBER: 32,327
; REFERENCE/DOCKET NUMBER: 218/078
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (213) 489-1600
; TELEFAX: (213) 955-0440
; TELEX: 67-3510
; INFORMATION FOR SEQ ID NO: 2710:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 54 base pairs
; TYPE: nucleic acid
```

```
;
; STRANDEDNESS: single
; TOPOLOGY: linear
US-09-038-073-2710

Query Match 100.0%; Score 10; DB 3; Length 54;
Best Local Similarity 100.0%; Pred. No. 1.9e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

RESULT 50
US-09-446-047A-1
; Sequence 1, Application US/09446047A
; Patent No. 6379924
; GENERAL INFORMATION:
; APPLICANT: Darrell Sleep
; APPLICANT: Delta Biotechnology Limited
; TITLE OF INVENTION: Improved Protein Expression Strains
; NUMBER OF SEQUENCES: 20
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Aventis Behring LLC
; STREET: 1020 First Avenue
; CITY: King of Prussia
; STATE: Pennsylvania
; COUNTRY: United States of America
; ZIP: 19406-1310
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30 (EPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/446,047A
; FILING DATE: 15-Dec-1999
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 70 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "PCR primer"
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-446-047A-1

Query Match 100.0%; Score 10; DB 4; Length 70;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 30 CTTCTCTTTT 39

RESULT 51
US-08-117-374-5
; Sequence 5, Application US/08117374
; Patent No. 5362865
; GENERAL INFORMATION:
; APPLICANT: Austin, Glenn D.
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; TITLE OF INVENTION: No. 5362865-translated Leader Sequences
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 5362865th
; CITY: St. Louis
; STATE: Missouri
```

; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: US/08/117,374
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10531)A
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 71 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-117-374-5

Query Match 100.0%; Score 10; DB 1; Length 71;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 43 CTTCTCTTTT 52

RESULT 52
US-08-280-263-5
; Sequence 5, Application US/08280263
; Patent No. 5659122
; GENERAL INFORMATION:
; APPLICANT: Austin, Glenn D.
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, B4F
; STREET: 700 Chesterfield Parkway No. 5659122th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; FILING DATE: 25-JUL-1994
; CLASSIFICATION: 800
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/117,374
; FILING DATE: 02-SEP-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10531) A
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:

; LENGTH: 71 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-280-263-5

Query Match 100.0%; Score 10; DB 1; Length 71;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 43 CTTCTCTTTT 52

RESULT 53
US-08-597-325-5
; Sequence 5, Application US/08597325
; Patent No. 6018100
; GENERAL INFORMATION:
; APPLICANT: Rogers, Stephen G.
; TITLE OF INVENTION: Promoter for Transgenic Plants
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, B4F
; STREET: 700 Chesterfield Parkway No. 6018100th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: 08/597,325
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/366,240
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10647)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 71 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; US-08-597-325-5

Query Match 100.0%; Score 10; DB 3; Length 71;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 43 CTTCTCTTTT 52

RESULT 54
US-08-597-325-5
; Sequence 5, Application US/08597325
; Patent No. 6018100
; GENERAL INFORMATION:

```

; APPLICANT: Rogers, Stephen G.
; TITLE OF INVENTION: Promoter for Transgenic Plants
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 6018100th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: 08/597,325
; APPLICATION NUMBER: US/08/597,325
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/366,240
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10647)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 71 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
;
US-08-597-325-5

Query Match 100.0%; Score 10; DB 3; Length 71;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 43 CTTCTCTTTT 52

RESULT 55
PCT-US94-10256-5
; Sequence 5, Application PCT/US9410256
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; TITLE OF INVENTION: Non-translated Leader Sequences
; NUMBER OF SEQUENCES: 22
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25 (EPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US94/10256
; FILING DATE: 01-SEPT-1994
; CLASSIFICATION:
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 71 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
;
PCT-US94-10256-5
```

```

Query Match 100.0%; Score 10; DB 5; Length 71;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 43 CTTCTCTTTT 52

RESULT 56
US-08-117-374-6/c
; Sequence 6, Application US/08117374
; Patent No. 5362865
; GENERAL INFORMATION:
; APPLICANT: Austin, Glenn D.
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; TITLE OF INVENTION: No. 5362865-translated Leader Sequences
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 5362865th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/117,374
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10531)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 75 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
;
US-08-117-374-6

Query Match 100.0%; Score 10; DB 1; Length 75;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 33 CTTCTCTTTT 24

RESULT 57
US-08-280-263-6/c
; Sequence 6, Application US/08280263
; Patent No. 5659122
; GENERAL INFORMATION:
; APPLICANT: Austin, Glenn D.
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; TITLE OF INVENTION: No. 5659122-translated Leader Sequences
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 5659122th
; CITY: St. Louis
; STATE: Missouri
```

```
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/280,263
; FILING DATE: 25-JUL-1994
; CLASSIFICATION: 800
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/117,374
; FILING DATE: 02-SEP-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10531)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 6:
; Sequence 6, Application US/08597325
; Patent No. 6018100
; GENERAL INFORMATION:
; APPLICANT: Rogers, Stephen G.
; TITLE OF INVENTION: Promoter for Transgenic Plants
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 6018100th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: 08/597,325
; APPLICATION NUMBER: US/08/597,325
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/366,240
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10647)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 6:
; Sequence 6, Application US/08597325
; Patent No. 6018100
; GENERAL INFORMATION:
; APPLICANT: Rogers, Stephen G.
; TITLE OF INVENTION: Promoter for Transgenic Plants
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 6018100th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: 08/597,325
; APPLICATION NUMBER: US/08/597,325
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/366,240
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10647)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
```

```
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 75 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-08-597-325-6

Query Match 100.0%; Score 10; DB 3; Length 75;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 33 CTTCTCTTTT 24

RESULT 59
US-08-597-325-6/c
; Sequence 6, Application US/08597325
; Patent No. 6018100
; GENERAL INFORMATION:
; APPLICANT: Rogers, Stephen G.
; TITLE OF INVENTION: Promoter for Transgenic Plants
; NUMBER OF SEQUENCES: 11
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 6018100th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA: 08/597,325
; APPLICATION NUMBER: US/08/597,325
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/366,240
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10647)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 75 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-08-597-325-6

Query Match 100.0%; Score 10; DB 3; Length 75;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 33 CTTCTCTTTT 24

RESULT 60
PCT-US94-10256-6/c
; Sequence 6, Application PC/TUS9410256
```

GENERAL INFORMATION:
; APPLICANT: Austin, Glenn D.
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; TITLE OF INVENTION: Non-translated Leader Sequences
; NUMBER OF SEQUENCES: 22
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25 (BPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US94/10256
; FILING DATE: 01-SEPT-1994
; CLASSIFICATION: 435
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 75 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
PCT-US94-10256-6

Query Match 100.0%; Score 10; DB 5; Length 75;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 33 CTTCTCTTTT 24

RESULT 61
US-08-117-374-19
; Sequence 19, Application US/08117374
; Patent No. 5362865
; GENERAL INFORMATION:
; APPLICANT: Austin, Glenn D.
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; TITLE OF INVENTION: Non-translated Leader Sequences
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 5362865th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/117,374
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10531)A
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 19:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 78 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-08-117-374-19

Query Match 100.0%; Score 10; DB 1; Length 78;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 43 CTTCTCTTTT 52

RESULT 62
US-08-280-263-19
; Sequence 19, Application US/08280263
; Patent No. 5659122
; GENERAL INFORMATION:
; APPLICANT: Austin, Glenn D.
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; TITLE OF INVENTION: Non-translated Leader Sequences
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle D. Strode, Monsanto Company, BB4F
; STREET: 700 Chesterfield Parkway No. 5659122th
; CITY: St. Louis
; STATE: Missouri
; COUNTRY: USA
; ZIP: 63198
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/280,263
; FILING DATE: 25-JUL-1994
; CLASSIFICATION: 800
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/117,374
; FILING DATE: 02-SEP-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Strode, Janelle D.
; REGISTRATION NUMBER: 34,738
; REFERENCE/DOCKET NUMBER: 38-21(10531)A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (314)537-6224
; TELEFAX: (314)537-6047
; INFORMATION FOR SEQ ID NO: 19:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 78 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
US-08-280-263-19

Query Match 100.0%; Score 10; DB 1; Length 78;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||
Db 43 CTTCTCTTTT 52

RESULT 63
PCT-US94-10256-19
; Sequence 19, Application PC/TUS9410256
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: Enhanced Expression in Plants Using
; TITLE OF INVENTION: Non-translated Leader Sequences
; NUMBER OF SEQUENCES: 22
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: PCT/US94/10256
 ; FILING DATE: 01-SEPT-1994
 ; CLASSIFICATION:
 ; INFORMATION FOR SEQ ID NO: 19:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 78 base pairs
 ; TYPE: nucleic acid
 ; STRANDEDNESS: double
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: DNA (genomic)
 ; PCT-US94-10256-19

Query Match 100.0%; Score 10; DB 5; Length 78;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 Db 43 CTTCTCTTTT 52

RESULT 64
 US-08-117-374-20/c
 ; Sequence 20, Application US/08117374
 ; Patent No. 5362865
 ; GENERAL INFORMATION:
 ; APPLICANT: Austin, Glenn D.
 ; TITLE OF INVENTION: Enhanced Expression in Plants Using
 ; TITLE OF INVENTION: No. 5362865-translated Leader Sequences
 ; NUMBER OF SEQUENCES: 22
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Janelle D. Strobe, Monsanto Company, BB4F
 ; STREET: 700 Chesterfield Parkway No. 5362865th
 ; CITY: St. Louis
 ; STATE: Missouri
 ; COUNTRY: USA
 ; ZIP: 63198
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: PatentIn Release #1.0, Version #1.25
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/117,374
 ; FILING DATE:
 ; CLASSIFICATION: 435
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Strobe, Janelle D.
 ; REGISTRATION NUMBER: 34,738
 ; REFERENCE/DOCKET NUMBER: 38-21(10531)A
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (314)537-6224
 ; TELEFAX: (314)537-6047
 ; INFORMATION FOR SEQ ID NO: 20:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 82 base pairs
 ; TYPE: nucleic acid
 ; STRANDEDNESS: double
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: DNA (genomic)
 ; US-08-117-374-20

Query Match 100.0%; Score 10; DB 1; Length 82;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 Db 40 CTTCTCTTTT 31

RESULT 65
 US-08-280-263-20/c
 ; Sequence 20, Application US/08280263
 ; Patent No. 5659122
 ; GENERAL INFORMATION:
 ; APPLICANT: Austin, Glenn D.
 ; TITLE OF INVENTION: Enhanced Expression in Plants Using
 ; TITLE OF INVENTION: No. 5659122-translated Leader Sequences
 ; NUMBER OF SEQUENCES: 22
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Janelle D. Strobe, Monsanto Company, BB4F
 ; STREET: 700 Chesterfield Parkway No. 5659122th
 ; CITY: St. Louis
 ; STATE: Missouri
 ; COUNTRY: USA
 ; ZIP: 63198
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: PatentIn Release #1.0, Version #1.25
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/08/280,263
 ; FILING DATE: 25-JUL-1994
 ; CLASSIFICATION: 800
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: US 08/117,374
 ; FILING DATE: 02-SEP-1993
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Strobe, Janelle D.
 ; REGISTRATION NUMBER: 34,738
 ; REFERENCE/DOCKET NUMBER: 38-21(10531)A
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: (314)537-6224
 ; TELEFAX: (314)537-6047
 ; INFORMATION FOR SEQ ID NO: 20:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 82 base pairs
 ; TYPE: nucleic acid
 ; STRANDEDNESS: double
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: DNA (genomic)
 ; US-08-280-263-20

Query Match 100.0%; Score 10; DB 1; Length 82;
 Best Local Similarity 100.0%; Pred. No. 1.8e+03;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
 Db 40 CTTCTCTTTT 31

RESULT 66
 PCT-US94-10256-20/c
 ; Sequence 20, Application PC/TUS9410256
 ; GENERAL INFORMATION:
 ; APPLICANT:
 ; TITLE OF INVENTION: Enhanced Expression in Plants Using
 ; TITLE OF INVENTION: Non-translated Leader Sequences
 ; NUMBER OF SEQUENCES: 22
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Floppy disk
 ; COMPUTER: IBM PC compatible
 ; OPERATING SYSTEM: PC-DOS/MS-DOS
 ; SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: PCT/US94/10256
 ; FILING DATE: 01-SEPT-1994
 ; CLASSIFICATION:
 ; INFORMATION FOR SEQ ID NO: 20:
 ; SEQUENCE CHARACTERISTICS:

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; LENGTH: 82 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
PCT-US94-10256-20

Query Match          100.0%; Score 10; DB 5; Length 82;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10
    |||||
Db 40 CTCTCTCTTT 31

RESULT 67
US-09-174-465D-11/c
; Sequence 11, Application US/09174465D
; Patent No. 6180364
; GENERAL INFORMATION:
; APPLICANT: KOMAN, Ahment
; APPLICANT: CHASSIN, Dorine
; APPLICANT: BELLET, Dominique
; TITLE OF INVENTION: NEW PROTEIN CALLED EPIL/PLACENTIN. PROCESS FOR THE
; PREPARATION OF THIS PROTEIN AND PHARMACEUTICAL
; TITLE OF INVENTION: PREPARATION OF THIS PROTEIN AND PHARMACEUTICAL
; TITLE OF INVENTION: COMPOSITION CONTAINING SUCH, DNA CODING FOR SAID
; TITLE OF INVENTION: PROTEIN
; FILE REFERENCE: 017753-103
; CURRENT APPLICATION NUMBER: US/09/174,465D
; CURRENT FILING DATE: 1998-10-19
; PRIOR APPLICATION NUMBER: US 08/482,842
; PRIOR FILING DATE: 1995-06-07
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 11
; LENGTH: 93
; TYPE: DNA
; ORGANISM: Unknown
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(93)
; OTHER INFORMATION: Description of Unknown Organism:EPIL - Early
; OTHER INFORMATION: Placenta Insulin-Like peptide
US-09-174-465D-11

Query Match          100.0%; Score 10; DB 3; Length 93;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10
    |||||
Db 17 CTCTCTCTTT 8

RESULT 68
US-09-599-564A-11/c
; Sequence 11, Application US/09599564A
; Patent No. 6362318
; GENERAL INFORMATION:
; APPLICANT: KOMAN, Ahment
; APPLICANT: CHASSIN, Dorine
; APPLICANT: BELLET, Dominique
; TITLE OF INVENTION: NEW PROTEIN CALLED EPIL/PLACENTIN. PROCESS FOR THE
; PREPARATION OF THIS PROTEIN AND PHARMACEUTICAL
; TITLE OF INVENTION: PREPARATION OF THIS PROTEIN AND PHARMACEUTICAL
; TITLE OF INVENTION: COMPOSITION CONTAINING SUCH, DNA CODING FOR SAID
; TITLE OF INVENTION: PROTEIN
; FILE REFERENCE: 017753-127
; CURRENT APPLICATION NUMBER: US/09/599,564A
; CURRENT FILING DATE: 2000-06-23
; PRIOR APPLICATION NUMBER: 09/174,465
; PRIOR FILING DATE: 1998-10-19
; PRIOR APPLICATION NUMBER: US 08/482,842

Query Match          100.0%; Score 10; DB 3; Length 93;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10
    |||||
Db 17 CTCTCTCTTT 8

RESULT 69
US-09-461-697-223/c
; Sequence 223, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Purnam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; TITLE OF INVENTION: CELL DEATH
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 223
; LENGTH: 96
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-223

Query Match          100.0%; Score 10; DB 3; Length 96;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTCTCTCTTT 10
    |||||
Db 68 CTCTCTCTTT 59

RESULT 70
US-09-144-428-55/c
; Sequence 55, Application US/09144428
; Patent No. 6583108
; GENERAL INFORMATION:
; APPLICANT: BAYER CORPORATION, The
; APPLICANT: TAMBURINI, Paul P
; APPLICANT: DAVIS, Gary
; APPLICANT: DELARIA, Katherine A
; APPLICANT: MARLOR, Christopher W
; APPLICANT: MULLER, Daniel K
; TITLE OF INVENTION: HUMAN BIKUNIN
; NUMBER OF SEQUENCES: 71
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: McDonnell Boehnen Hulbert & Berghoff
```

; STREET: 300 S. Wacker Drive Suite 3200
; CITY: CHICAGO
; STATE: ILLINOIS
; COUNTRY: USA
; ZIP: 60606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/144.428
; FILING DATE: 10-MAR-1997
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/US97/03894
; FILING DATE: 10-MAR-1997
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/013.106
; FILING DATE: 11-MAR-1996
; APPLICATION DATA:
; APPLICATION NUMBER: US 60/019.793
; FILING DATE: 14-JUN-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/725.251
; FILING DATE: 04-OCT-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: CHAO, Mark
; REGISTRATION NUMBER: 37,293
; REFERENCE/DOCKET NUMBER: 96,223-II
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (312) 913-0001
; TELEFAX: (312) 913-0002
; INFORMATION FOR SEQ ID NO: 55:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 102 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: unknown
; TOPOLOGY: unknown
; MOLECULE TYPE: DNA (genomic)
; US-09-144-428-55

Query Match 100.0%; Score 10; DB 4; Length 102;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||
Db 28 CTTCTCTTTT 19

RESULT 71
US-08-932-082-10
; Sequence 10, Application US/08932082
; Patent No. 6251856
; GENERAL INFORMATION:
; APPLICANT: Markussen, Jan
; APPLICANT: Jonassen, Ib
; APPLICANT: Havelund, Svend
; APPLICANT: Brandt, Jakob
; APPLICANT: Kurtzhals, Peter
; APPLICANT: Hansen, Hertz Per
; APPLICANT: Kaarsholm, Niels Christian
; TITLE OF INVENTION: INSULIN DERIVATIVES
; NUMBER OF SEQUENCES: 26
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: No. 6251856 No. 6251856disk of No. 6251856th America, Inc.
; STREET: 405 Lexington Avenue, 64th Floor
; CITY: New York
; STATE: New York
; COUNTRY: United States of America
; ZIP: 10174-6401

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/932,082
; FILING DATE: 12-AUG-1997
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Lambiris, Elias J.
; REGISTRATION NUMBER: 33,728
; REFERENCE/DOCKET NUMBER: 4341.204-US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 212-867-0123
; TELEFAX: 212-878-9655
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 112 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; US-08-932-082-10

Query Match 100.0%; Score 10; DB 3; Length 112;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||
Db 88 CTTCTCTTTT 97

RESULT 72

US-09-461-697-221/c
; Sequence 221, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-085-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 221
; LENGTH: 117
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-461-697-221

Query Match 100.0%; Score 10; DB 3; Length 117;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||
Db 89 CTTCTCTTTT 80

RESULT 73

US-09-461-697-219/c
; Sequence 219, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:

```
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 219
; LENGTH: 126
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-219

Query Match      100.0%; Score 10; DB 3; Length 126;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      98 CTTCTCTTTT 89
|||||

RESULT 74
US-09-441-416A-22/c
; Sequence 22, Application US/09441416A
; Patent No. 6294518
; GENERAL INFORMATION:
; APPLICANT: Potter, David A.
; APPLICANT: Skolnik, Paul R.
; TITLE OF INVENTION: CELL-PERMEABLE PROTEIN INHIBITORS OF
; FILE REFERENCE: 00398-140001
; CURRENT APPLICATION NUMBER: US/09/441,416A
; CURRENT FILING DATE: 1999-11-16
; PRIOR APPLICATION NUMBER: US 08/964,302
; PRIOR FILING DATE: 1997-11-04
; NUMBER OF SEQ ID NOS: 23
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 22
; LENGTH: 132
; TYPE: DNA
; ORGANISM: Eukaryote
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (7)...(129)
US-09-441-416A-22

Query Match      100.0%; Score 10; DB 3; Length 132;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      79 CTTCTCTTTT 70
|||||

RESULT 75
US-08-477-928A-36/c
; Sequence 36, Application US/08477928A
; Patent No. 6207389
; GENERAL INFORMATION:
; APPLICANT: Dosch, Hans M.
; TITLE OF INVENTION: METHODS FOR CONTROLLING T
; TITLE OF INVENTION: LYMPHOCYTE MEDIATED IMMUNE RESPONSES
; NUMBER OF SEQUENCES: 49
```

```
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BAKER & BOTTS
; STREET: 1299 Pennsylvania Avenue
; CITY: Washington D.C.
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 20004-2400
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/477,928A
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 536
; ATTORNEY/AGENT INFORMATION:
; NAME: Remenick, James
; REGISTRATION NUMBER: 36902
; REFERENCE/DOCKET NUMBER: 19060-0105
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 639 7700
; TELEFAX: (202) 639 7890
; INFORMATION FOR SEQ ID NO: 36:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 146 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-477-928A-36

Query Match      100.0%; Score 10; DB 3; Length 146;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      138 CTTCTCTTTT 129
|||||

RESULT 76
US-09-461-697-217/c
; Sequence 217, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 217
; LENGTH: 156
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-217

Query Match      100.0%; Score 10; DB 3; Length 156;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      138 CTTCTCTTTT 129
|||||
```

Db 128 CTTCTCTTTT 119

RESULT 77

US-09-461-697-215/c
; Sequence 215, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461.697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 215
; LENGTH: 174
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-215

Query Match 100.0%; Score 10; DB 3; Length 174;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 146 CTTCTCTTTT 137

RESULT 78

US-09-461-697-213/c
; Sequence 213, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461.697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 213
; LENGTH: 189
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-213

Query Match 100.0%; Score 10; DB 3; Length 189;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||

Db 161 CTTCTCTTTT 152

RESULT 79

US-09-461-697-211/c
; Sequence 211, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461.697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 211
; LENGTH: 195
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-211

Query Match 100.0%; Score 10; DB 3; Length 195;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 167 CTTCTCTTTT 158

RESULT 80

US-09-107-532A-2918/c
; Sequence 2918, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Walcham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD-ROM ISO9660
; COMPUTER: PC
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107.532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 2918:
; SEQUENCE CHARACTERISTICS:

```
;
; LENGTH: 198 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Enterococcus faecium
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (B) LOCATION 1...198
; SEQUENCE DESCRIPTION: SEQ ID NO: 2918:
US-09-107-532A-2918

Query Match      100.0%; Score 10; DB 4; Length 198;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      126 CTTCTCTTTT 117

RESULT 81
US-09-107-532A-439
; Sequence 439, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD-ROM ISO9660
; COMPUTER: PC
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 439:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 201 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Enterococcus faecium
; FEATURE:
; NAME/KEY: misc_feature
```

```
;
; LOCATION: (B) LOCATION 1...201
; SEQUENCE DESCRIPTION: SEQ ID NO: 439:
US-09-107-532A-439

Query Match      100.0%; Score 10; DB 4; Length 201;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      177 CTTCTCTTTT 186

RESULT 82
US-09-107-532A-3253/c
; Sequence 3253, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD-ROM ISO9660
; COMPUTER: PC
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 3253:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 201 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Enterococcus faecium
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (B) LOCATION 1...201
; SEQUENCE DESCRIPTION: SEQ ID NO: 3253:
US-09-107-532A-3253

Query Match      100.0%; Score 10; DB 4; Length 201;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      178 CTTCTCTTTT 169
```

RESULT 83
US-09-313-294A-7004
; Sequence 7004, Application US/09313294A
; Patent No. 6476212
; GENERAL INFORMATION:
; APPLICANT: Lalgudi, Raghunath V.
; APPLICANT: Ito, Laura Y.
; APPLICANT: Sherman, Bradley K.
; TITLE OF INVENTION: POLYNUCLEOTIDES AND POLYPEPTIDES DERIVED FROM CORN EAR
; FILE REFERENCE: PL-0017 US
; CURRENT APPLICATION NUMBER: US/09/313,294A
; CURRENT FILING DATE: 1999-05-14
; NUMBER OF SEQ ID NOS: 7600
; SOFTWARE: PERL Program
; SEQ ID NO 7004
; LENGTH: 206
; TYPE: DNA
; ORGANISM: Zea mays
; FEATURE:
; NAME/KEY: misc.feature
; OTHER INFORMATION: Incyte ID No. 6476212 700380982H1
; NAME/KEY: unsure
; LOCATION: 190, 193
; OTHER INFORMATION: a, t, c, g, or other
US-09-313-294A-7004

Query Match 100.0%; Score 10; DB 4; Length 206;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
DB 129 CTTCTCTTTT 138

RESULT 84
US-09-461-697-209/c
; Sequence 209, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 209
; LENGTH: 213
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-209

Query Match 100.0%; Score 10; DB 3; Length 213;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
DB 185 CTTCTCTTTT 176

RESULT 85

5217891-19/c
; Patent No. 5217891
; APPLICANT: BRAKE, ANTHONY J.; VAN DEN BERG, JOHAN A.
; TITLE OF INVENTION: DNA CONSTRUCTS CONTAINING A KLUYVEROMYCES
; A FACTOR LEADER SEQUENCE FOR DIRECTING SECRETION OF HETEROLOGOUS
; POLYPEPTIDES
; NUMBER OF SEQUENCES: 23
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/507,398
; FILING DATE: 09-APR-1990
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 78,551
; FILING DATE: 28-JUL-1987
; SEQ ID NO:19:
; LENGTH: 222
5217891-19
Query Match 100.0%; Score 10; DB 6; Length 222;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
DB 37 CTTCTCTTTT 28
RESULT 86
US-09-016-434-99/c
; Sequence 99, Application US/09016434
; Patent No. 6500938
; GENERAL INFORMATION:
; APPLICANT: Janice Au-Young
; APPLICANT: Jeffrey J. Seilhamer
; TITLE OF INVENTION: COMPOSITION FOR THE DETECTION OF SIGNALING
; TITLE OF INVENTION: PATHWAY GENE EXPRESSION
; NUMBER OF SEQUENCES: 1490
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: INCYTE PHARMACEUTICALS, INC.
; STREET: 3174 PORTER DRIVE
; CITY: PALO ALTO
; STATE: CALIFORNIA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Word Perfect 6.1 for Windows/MS-DOS 6.2
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/016,434
; FILING DATE: HERewith
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Zeller, Karen J.
; REGISTRATION NUMBER: 37,071
; REFERENCE/DOCKET NUMBER: PA-0002 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650) 855-0555
; TELEFAX: (650) 845-4166
; INFORMATION FOR SEQ ID NO: 99:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 228 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: LUNGFET03
; CLONE: 1251228
US-09-016-434-99

```
Query Match      100.0%; Score 10; DB 4; Length 228;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTCTCTCTTT 10
DB      58 CTCTCTCTTT 49

RESULT 87
US-09-461-697-207/c
; Sequence 207, Application US/09461697
; Patent No. 6277974
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/461,697
; CURRENT FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 207
; LENGTH: 231
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-461-697-207

Query Match      100.0%; Score 10; DB 3; Length 231;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTCTCTCTTT 10
DB      203 CTCTCTCTTT 194

RESULT 88
US-09-107-532A-2834/c
; Sequence 2834, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD-ROM ISO9660
; COMPUTER: PC
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
```

```
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 2834:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 231 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Enterococcus faecium
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (B) LOCATION 1...231
; SEQUENCE DESCRIPTION: SEQ ID NO: 2834:
US-09-107-532A-2834
; Query Match      100.0%; Score 10; DB 4; Length 231;
; Best Local Similarity 100.0%; Pred. No. 1.8e+03;
; Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTCTCTCTTT 10
DB      79 CTCTCTCTTT 70

RESULT 89
US-09-107-532A-724/c
; Sequence 724, Application US/09107532A
; Patent No. 6583275
; GENERAL INFORMATION:
; APPLICANT: Lynn A Doucette-Stamm and David Bush
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
; NUMBER OF SEQUENCES: 7310
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: GENOME THERAPEUTICS CORPORATION
; STREET: 100 Beaver Street
; CITY: Waltham
; STATE: Massachusetts
; COUNTRY: USA
; ZIP: 02354
; COMPUTER READABLE FORM:
; MEDIUM TYPE: CD-ROM ISO9660
; COMPUTER: PC
; OPERATING SYSTEM: <Unknown>
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/107,532A
; FILING DATE: 30-Jun-1998
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/085,598
; FILING DATE: 14 May 1998
; APPLICATION NUMBER: 60/051571
; FILING DATE: July 2, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Ariniello, Pamela Deneke
; REGISTRATION NUMBER: 40,489
; REFERENCE/DOCKET NUMBER: GTC-012
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (781)893-5007
; TELEFAX: (781)893-8277
; INFORMATION FOR SEQ ID NO: 724:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 234 base pairs
```



```

; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Enterococcus faecium
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (8) LOCATION 1...234
; SEQUENCE DESCRIPTION: SEQ ID NO: 724:
US-09-107-532A-724

Query Match 100.0%; Score 10; DB 4; Length 234;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 118 CTTCTCTTTT 109

RESULT 90
US-09-397-787-56
; Sequence 56, Application US/09397787
; Patent No. 6468758
; GENERAL INFORMATION:
; APPLICANT: Benson, Darin R.
; APPLICANT: Lodes, Michael J.
; APPLICANT: Mitcham, Jennifer L.
; APPLICANT: King, Gordon E.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR OVARIAN
; TITLE OF INVENTION: CANCER THERAPY AND DIAGNOSIS
; FILE REFERENCE: 210121.466C2
; CURRENT APPLICATION NUMBER: US/09/397,787
; CURRENT FILING DATE: 1999-09-16
; NUMBER OF SEQ ID NOS: 334
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 56
; LENGTH: 241
; TYPE: DNA
; ORGANISM: Homo sapien
US-09-397-787-56

Query Match 100.0%; Score 10; DB 4; Length 241;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 71 CTTCTCTTTT 80

RESULT 91
US-09-389-681-360
; Sequence 360, Application US/09389681A
; Patent No. 6518237
; GENERAL INFORMATION:
; APPLICANT: Yuqui, Jiang
; APPLICANT: Dillon, Davin C.
; APPLICANT: Mitcham, Jennifer L.
; APPLICANT: Xu, Jiangchun
; APPLICANT: Harlocker, Susan L.
; TITLE OF INVENTION: COMPOSITIONS OF BREAST CANCER AND METHODS FOR THEIR USE
; FILE REFERENCE: 210121.470C3
; CURRENT APPLICATION NUMBER: US/09/389,681A
; CURRENT FILING DATE: 1999-09-02
; NUMBER OF SEQ ID NOS: 463
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 360
; LENGTH: 241
; TYPE: DNA

; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: circular
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)...(241)
; OTHER INFORMATION: n = A,T,C or G
US-09-389-681-360

Query Match 100.0%; Score 10; DB 4; Length 241;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 113 CTTCTCTTTT 122

RESULT 92
US-09-620-405B-360
; Sequence 360, Application US/09620405B
; Patent No. 6528054
; GENERAL INFORMATION:
; APPLICANT: Jiang, Yuqui
; APPLICANT: Dillon, Davin C.
; APPLICANT: Mitcham, Jennifer L.
; APPLICANT: Xu, Jiangchun
; APPLICANT: Harlocker, Susan L.
; APPLICANT: Hepler, William T.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND
; FILE REFERENCE: 210121.470C8
; CURRENT APPLICATION NUMBER: US/09/620,405B
; CURRENT FILING DATE: 2000-07-20
; NUMBER OF SEQ ID NOS: 495
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 360
; LENGTH: 241
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)...(241)
; OTHER INFORMATION: n = A,T,C or G
US-09-620-405B-360

Query Match 100.0%; Score 10; DB 4; Length 241;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 113 CTTCTCTTTT 122

RESULT 93
US-09-433-826B-360
; Sequence 360, Application US/09433826B
; Patent No. 6579973
; GENERAL INFORMATION:
; APPLICANT: Jiang, Yuqui
; APPLICANT: Dillon, Davin C.
; APPLICANT: Mitcham, Jennifer L.
; APPLICANT: Xu, Jiangchun
; APPLICANT: Harlocker, Susan L.
; TITLE OF INVENTION: COMPOSITIONS OF BREAST CANCER AND METHODS FOR THEIR USE
; FILE REFERENCE: 210121.470C4
; CURRENT APPLICATION NUMBER: US/09/433,826B
; CURRENT FILING DATE: 1999-11-03
; NUMBER OF SEQ ID NOS: 474
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 360
; LENGTH: 241
; TYPE: DNA
```

```

; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(241)
; OTHER INFORMATION: n = A,T,C or G
US-09-433-826B-360

Query Match      100.0%; Score 10; DB 4; Length 241;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      113 CTTCTCTTTT 122

RESULT 94
US-09-604-287A-360
; Sequence 360, Application US/09604287A
; Patent No. 6586572
; GENERAL INFORMATION:
; APPLICANT: Jiang, Yugu
; APPLICANT: Dillon, Devin C.
; APPLICANT: Mitcham, Jennifer L.
; APPLICANT: Xu, Jiangchun
; APPLICANT: Harlocker, Susan L.
; APPLICANT: Hepler, William T.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND
; TITLE OF INVENTION: DIAGNOSIS OF BREAST CANCER
; FILE REFERENCE: 210121.470C7
; CURRENT APPLICATION NUMBER: US/09/604,287A
; CURRENT FILING DATE: 2000-06-22
; NUMBER OF SEQ ID NOS: 489
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 360
; LENGTH: 241
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(241)
; OTHER INFORMATION: n = A,T,C or G
US-09-604-287A-360

Query Match      100.0%; Score 10; DB 4; Length 241;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      113 CTTCTCTTTT 122

RESULT 95
US-09-016-434-737/c
; Sequence 737, Application US/09016434
; Patent No. 6500938
; GENERAL INFORMATION:
; APPLICANT: Janice Au-Young
; APPLICANT: Jeffrey J. Seilhamer
; TITLE OF INVENTION: COMPOSITION FOR THE DETECTION OF SIGNALING
; TITLE OF INVENTION: PATHWAY GENE EXPRESSION
; NUMBER OF SEQUENCES: 1490
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: INCYTE PHARMACEUTICALS, INC.
; STREET: 3174 PORTER DRIVE
; CITY: PALO ALTO
; STATE: CALIFORNIA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Word Perfect 6.1 for Windows/MS-DOS 6.2
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/098,789
; FILING DATE: Herewith
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: CERRONE, MICHAEL C.
; REGISTRATION NUMBER: 39,132
; REFERENCE/DOCKET NUMBER: PF-0547 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650) 855-0555
; TELEFAX: (650) 845-4166

```

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; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Word Perfect 6.1 for Windows/MS-DOS 6.2
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/016,434
; FILING DATE: HEREWITH
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Zeller, Karen J.
; REGISTRATION NUMBER: 37,071
; REFERENCE/DOCKET NUMBER: PA-0002 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650) 855-0555
; TELEFAX: (650) 845-4166
; INFORMATION FOR SEQ ID NO: 737:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 247 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: TMLR2DT01
; CLONE: 395476
US-09-016-434-737

Query Match      100.0%; Score 10; DB 4; Length 247;
Best Local Similarity 100.0%; Pred. No. 1.8e+03;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      118 CTTCTCTTTT 109

RESULT 96
US-09-098-789-9
; Sequence 9, Application US/09098789
; Patent No. 6180342
; GENERAL INFORMATION:
; APPLICANT: Bandman, Olga
; APPLICANT: Tang, Y. Tom
; APPLICANT: Lal, Preeti
; APPLICANT: Corley, Neil C.
; APPLICANT: Guegler, Karl J.
; APPLICANT: Patterson, Chandra
; TITLE OF INVENTION: VACULAR PROTON ATPASE SUBUNITS
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: INCYTE PHARMACEUTICALS, INC.
; STREET: 3174 PORTER DRIVE
; CITY: PALO ALTO
; STATE: CALIFORNIA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Word Perfect 6.1 for Windows/MS-DOS 6.2
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/098,789
; FILING DATE: Herewith
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: CERRONE, MICHAEL C.
; REGISTRATION NUMBER: 39,132
; REFERENCE/DOCKET NUMBER: PF-0547 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650) 855-0555
; TELEFAX: (650) 845-4166

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/ INFORMATION FOR SEQ ID NO: 9:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 251 base pairs
/ TYPE: nucleic acid
/ STRANDEDNESS: single
/ TOPOLOGY: linear
/ IMMEDIATE SOURCE:
/ LIBRARY: HIPONON02
/ CLONE: 2246348CT1
US-09-098-789-9

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/ Sequence 1077, Application US/09702705
/ Patent No. 6504010
/ GENERAL INFORMATION:
/ APPLICANT: Wang, Tongtong
/ APPLICANT: Bangur, Chaitanya S.
/ APPLICANT: Lodes, Michael A.
/ APPLICANT: Fanger, Gary
/ APPLICANT: Vedvick, Tom
/ APPLICANT: Carter, Darrick
/ APPLICANT: Retter, Marc
/ APPLICANT: Mannion, Jane
/ APPLICANT: Fan, Liqun
/ TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND
/ FILE REFERENCE: 210121.478C14
/ CURRENT APPLICATION NUMBER: US/09/702,705
/ CURRENT FILING DATE: 2000-10-30
/ NUMBER OF SEQ ID NOS: 1833
/ SOFTWARE: FastSeq for Windows Version 3.0
/ SEQ ID NO 1077
/ LENGTH: 256
/ TYPE: DNA
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US-09-702-705-1077

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/ Sequence 1077, Application US/09736457
/ Patent No. 6509448
/ GENERAL INFORMATION:
/ APPLICANT: Wang, Tongtong
/ APPLICANT: Bangur, Chaitanya S.
/ APPLICANT: Lodes, Michael A.
/ APPLICANT: Fanger, Gary
/ APPLICANT: Vedvick, Tom
/ APPLICANT: Carter, Darrick
/ APPLICANT: Retter, Marc
/ APPLICANT: Mannion, Jane
/ APPLICANT: Fan, Liqun
/ APPLICANT: Wang, Aijun
/ TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND

/ FILE REFERENCE: 210121.478C15
/ CURRENT APPLICATION NUMBER: US/09/736,457
/ CURRENT FILING DATE: 2000-12-13
/ NUMBER OF SEQ ID NOS: 1864
/ SOFTWARE: FastSeq for Windows Version 3.0
/ SEQ ID NO 1077
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/ ORGANISM: Homo sapien
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/ Sequence 3143, Application US/09313294A
/ Patent No. 6476212
/ GENERAL INFORMATION:
/ APPLICANT: Lalgudi, Raghunath V.
/ APPLICANT: Ito, Laura Y.
/ APPLICANT: Sherman, Bradley K.
/ TITLE OF INVENTION: POLYNUCLEOTIDES AND POLYPEPTIDES DERIVED FROM CORN EAR
/ FILE REFERENCE: PL-0017 US
/ CURRENT APPLICATION NUMBER: US/09/313,294A
/ CURRENT FILING DATE: 1999-05-14
/ NUMBER OF SEQ ID NOS: 7600
/ SOFTWARE: PERL Program
/ SEQ ID NO 3143
/ LENGTH: 263
/ TYPE: DNA
/ ORGANISM: Zea mays
/ NAME/KEY: misc feature
/ OTHER INFORMATION: Incyte ID No. 6476212 700610958H1
/ NAME/KEY: unsure
/ LOCATION: 216, 246
/ OTHER INFORMATION: a, t, c, g, or other
US-09-313-294A-3143

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/ Sequence 18, Application US/09134001C
/ Patent No. 6380370
/ GENERAL INFORMATION:
/ APPLICANT: Lynn Doucette-Stamm et al
/ TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO STAPHYLOCOCCUS
/ FILE REFERENCE: GTC-007
/ CURRENT APPLICATION NUMBER: US/09/134,001C
/ CURRENT FILING DATE: 1998-08-13
/ PRIOR APPLICATION NUMBER: US 60/064,964
/ PRIOR FILING DATE: 1997-11-08
/ PRIOR APPLICATION NUMBER: US 60/055,779
/ PRIOR FILING DATE: 1997-08-14
/ NUMBER OF SEQ ID NOS: 5674
/ SEQ ID NO 18
/ LENGTH: 267

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US-09-134-001C-18

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152	10	100.0	163	9	US-09-770-696-391	Sequence 391, App	225	10	100.0	250	9	US-09-880-107-2807	Sequence 2807, App
153	10	100.0	163	14	US-10-060-036-2004	Sequence 2004, App	226	10	100.0	253	12	US-09-237-183A-2137	Sequence 2137, App
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157	10	100.0	173	9	US-09-770-696-331	Sequence 331, App	230	10	100.0	256	10	US-09-849-626-1077	Sequence 1077, App
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159	10	100.0	174	10	US-09-922-261-215	Sequence 215, App	232	10	100.0	256	11	US-10-113-872-1077	Sequence 1077, App
160	10	100.0	175	12	US-09-814-353-6379	Sequence 6379, App	233	10	100.0	256	12	US-10-017-754-1077	Sequence 1077, App
161	10	100.0	175	12	US-09-814-353-12656	Sequence 12656, A	234	10	100.0	256	14	US-10-017-754-1077	Sequence 1077, App
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C 289	10	100.0	277	10	US-09-878-574-12185	Sequence 12185, A	C 362	10	100.0	316	10	US-09-736-457-1553	Sequence 1553, App
C 290	10	100.0	278	9	US-09-294-093B-5812	Sequence 5812, App	C 363	10	100.0	316	10	US-09-814-353-1535	Sequence 1553, App
C 291	10	100.0	279	10	US-09-878-574-10946	Sequence 10946, A	C 364	10	100.0	316	11	US-09-849-626-1553	Sequence 1553, App
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C 433	10	100.0	336	10	US-09-974-300-8266	Sequence 8266, Ap							
C 434	10	100.0	336	12	US-10-238-075-555	Sequence 555, App							
C 435	10	100.0	337	10	US-09-929-493-7	Sequence 7, Appli							
C 436	10	100.0	337	11	US-09-803-719-1668	Sequence 1668, Ap							
C 437	10	100.0	337	11	US-09-918-995-19793	Sequence 19793, A							
C 438	10	100.0	338	12	US-10-270-487-7	Sequence 7, Appli							
C 439	10	100.0	338	11	US-09-803-719-963	Sequence 963, App							
C 440	10	100.0	339	13	US-10-027-632-128730	Sequence 128730							
C 441	10	100.0	339	14	US-10-060-036-1555	Sequence 1555, Ap							
C 442	10	100.0	339	14	US-10-313-542-168	Sequence 168, App							
C 443	10	100.0	340	10	US-09-920-300A-423	Sequence 423, App							
C 444	10	100.0	340	12	US-10-099-926-423	Sequence 423, App							
C 445	10	100.0	340	13	US-10-033-528-423	Sequence 423, App							
C 446	10	100.0	342	9	US-09-770-791-814	Sequence 814, App							
C 447	10	100.0	342	11	US-09-918-995-18543	Sequence 18543, A							
C 448	10	100.0	342	12	US-10-116-712-589	Sequence 589, App							
C 449	10	100.0	343	11	US-09-803-719-1667	Sequence 1667, Ap							
C 450	10	100.0	343	12	US-10-125-159-81	Sequence 81, Appl							
C 451	10	100.0	344	12	US-10-116-712-134	Sequence 134, App							
C 452	10	100.0	345	9	US-09-770-791-775	Sequence 775, App							
C 453	10	100.0	346	10	US-09-969-708-371	Sequence 371, App							
C 454	10	100.0	346	10	US-09-969-708-436	Sequence 436, App							

ALIGNMENTS

RESULT 1

US-09-827-998-287/c
; Sequence 287, Application US/09827998
; Patent No. US20020102252A1

; GENERAL INFORMATION:

; APPLICANT: Gu, Yizhong

; APPLICANT: Shannon, Mark

; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E

; FILE REFERENCE: MDHWORF-8

; CURRENT APPLICATION NUMBER: US/09/827,998

; CURRENT FILING DATE: 2001-04-06

; PRIOR APPLICATION NUMBER: US 60/207,456

; PRIOR FILING DATE: 2000-05-26

; PRIOR APPLICATION NUMBER: US 60/236,359

; PRIOR FILING DATE: 2000-09-27

; NUMBER OF SEQ ID NOS: 1881

; SOFTWARE: Aecomica Sequence Listing Engine

; SEQ ID NO 287

; LENGTH: 17

; TYPE: DNA

; ORGANISM: Homo sapiens

US-09-827-998-287

Query Match 100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
| | | | | | | |
Db 17 CTTCTCTTTT 8

RESULT 2
US-09-827-998-288/c
; Sequence 288, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHWORF-8
; CURRENT APPLICATION NUMBER: US 60/207,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 288
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-288

Query Match 100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
| | | | | | | |
Db 16 CTTCTCTTTT 7

RESULT 3
US-09-827-998-289/c
; Sequence 289, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHWORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 289
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-289

Query Match 100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
| | | | | | | |
Db 15 CTTCTCTTTT 6

RESULT 4
US-09-827-998-290/c
; Sequence 290, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHWORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 290
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-290

Query Match 100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
| | | | | | | |
Db 14 CTTCTCTTTT 5

RESULT 5
US-09-827-998-291/c
; Sequence 291, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHWORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 291
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-291

Query Match 100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
| | | | | | | |
Db 13 CTTCTCTTTT 4

RESULT 6
US-09-827-998-292/c
; Sequence 292, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E

; FILE REFERENCE: MDHMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 292
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-292

Query Match 100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 12 CTTCTCTTTT 3

RESULT 7

US-09-827-998-293/c
; Sequence 293, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 293
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-293

Query Match 100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 11 CTTCTCTTTT 2

RESULT 8

US-09-827-998-294/c
; Sequence 294, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDHMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine

; SEQ ID NO 294
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-294

Query Match 100.0%; Score 10; DB 10; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 10 CTTCTCTTTT 1

RESULT 9

US-09-930-423-1740/c
; Sequence 1740, Application US/09930423
; Publication No. US20030092003A1
; GENERAL INFORMATION:
; APPLICANT: Blatt, Larry
; APPLICANT: McSwiggen, Jim
; TITLE OF INVENTION: Method and Reagent for the Treatment of Alzheimer's Disease
; FILE REFERENCE: MEB00,918-A 400/027
; CURRENT APPLICATION NUMBER: US/09/930,423
; CURRENT FILING DATE: 2001-08-15
; NUMBER OF SEQ ID NOS: 4553
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1740
; LENGTH: 17
; TYPE: RNA
; ORGANISM: Homo Sapiens
US-09-930-423-1740

Query Match 100.0%; Score 10; DB 11; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 14 CTTCTCTTTT 5

RESULT 10

US-09-930-423-1741/c
; Sequence 1741, Application US/09930423
; Publication No. US20030092003A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: Blatt, Larry
; APPLICANT: McSwiggen, Jim
; TITLE OF INVENTION: Method and Reagent for the Treatment of Alzheimer's Disease
; FILE REFERENCE: MEB00,918-A 400/027
; CURRENT APPLICATION NUMBER: US/09/930,423
; CURRENT FILING DATE: 2001-08-15
; NUMBER OF SEQ ID NOS: 4553
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1741
; LENGTH: 17
; TYPE: RNA
; ORGANISM: Homo Sapiens
US-09-930-423-1741

Query Match 100.0%; Score 10; DB 11; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 12 CTTCTCTTTT 3

```
RESULT 11
US-09-745-237A-1740/c
; Sequence 1740, Application US/09745237A
; Publication No. US20030143708A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: Blatt, Larry
; APPLICANT: McSwiggen, Jim
; TITLE OF INVENTION: Method and Reagent for the Treatment of Alzheimer's Disease
; FILE REFERENCE: 400/007 (MEHB00-918-A)
; CURRENT APPLICATION NUMBER: US/09/745,237A
; CURRENT FILING DATE: 2002-04-15
; NUMBER OF SEQ ID NOS: 4550
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1740
; LENGTH: 17
; TYPE: RNA
; ORGANISM: Homo sapiens
US-09-745-237A-1740

Query Match          100.0%; Score 10; DB 12; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      14 CTTCTCTTTT 5

RESULT 12
US-09-745-237A-1741/c
; Sequence 1741, Application US/09745237A
; Publication No. US20030143708A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: Blatt, Larry
; APPLICANT: McSwiggen, Jim
; TITLE OF INVENTION: Method and Reagent for the Treatment of Alzheimer's Disease
; FILE REFERENCE: 400/007 (MEHB00-918-A)
; CURRENT APPLICATION NUMBER: US/09/745,237A
; CURRENT FILING DATE: 2002-04-15
; NUMBER OF SEQ ID NOS: 4550
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 1741
; LENGTH: 17
; TYPE: RNA
; ORGANISM: Homo sapiens
US-09-745-237A-1741

Query Match          100.0%; Score 10; DB 12; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      12 CTTCTCTTTT 3

RESULT 13
US-10-060-756A-1720/c
; Sequence 1720, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 1720
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1720/c

Query Match          100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      16 CTTCTCTTTT 7

RESULT 14
US-10-060-756A-1721/c
; Sequence 1721, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US/09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US/60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 1721
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1721

Query Match          100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
Db      17 CTTCTCTTTT 8

RESULT 15
US-10-060-756A-1722/c
```

```
; Sequence 1722, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1722
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1722

Query Match      100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      15 CTTCTCTTTT 6

RESULT 16
US-10-060-756A-1723/c
; Sequence 1723, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1723
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
```

```
US-10-060-756A-1723
Query Match      100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      14 CTTCTCTTTT 5

RESULT 17
US-10-060-756A-1724/c
; Sequence 1724, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1724
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1724

Query Match      100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 CTTCTCTTTT 10
Db      13 CTTCTCTTTT 4

RESULT 18
US-10-060-756A-1725/c
; Sequence 1725, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
```

;
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1725
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1725

Query Match 100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||
Db 12 CTTCTCTTTT 3

RESULT 19
US-10-060-756A-1725/c
; Sequence 1726, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1726
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1726

Query Match 100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||
Db 11 CTTCTCTTTT 2

RESULT 20
US-10-060-756A-1727/c
; Sequence 1727, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian

;
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1727
; LENGTH: 17
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-1727

Query Match 100.0%; Score 10; DB 14; Length 17;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||
Db 10 CTTCTCTTTT 1

RESULT 21
US-10-205-309-284/c
; Sequence 284, Application US/10205309
; Publication No. US20030190635A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: McSwiggen, James
; TITLE OF INVENTION: RNA Interference Mediated Inhibition of Alzheimer's Disease Usin
; FILE REFERENCE: 900/033
; CURRENT APPLICATION NUMBER: US/10/205,309
; CURRENT FILING DATE: 2002-10-25
; NUMBER OF SEQ ID NOS: 674
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 284
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Target sequence/siNA sense
US-10-205-309-284

Query Match 100.0%; Score 10; DB 12; Length 19;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||
Db 10 CTTCTCTTTT 1

RESULT 22
US-10-205-309-609
; Sequence 609, Application US/10205309
; Publication No. US20030190635A1
; GENERAL INFORMATION:

```
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: McSwiggen, James
; TITLE OF INVENTION: RNA Interference Mediated Inhibition of Alzheimer's Disease Using
; TITLE OF INVENTION: Interfering RNA
; FILE REFERENCE: 900/033
; CURRENT APPLICATION NUMBER: US/10/205,309
; CURRENT FILING DATE: 2002-10-25
; NUMBER OF SEQ ID NOS: 674
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 609
; LENGTH: 19
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: siNA antisense region
US-10-205-309-609

Query Match      100.0%; Score 10; DB 12; Length 19;
Best Local Similarity 30.0%; Pred. No. 1.6e+04;
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
   |:|:|:|:|
Db 10 CUUCUCUUU 19

RESULT 23
US-10-087-325-15/c
; Sequence 15, Application US/10087325
; Publication No. US20020192682A1
; GENERAL INFORMATION:
; APPLICANT: Escary, Jean-Louis
; TITLE OF INVENTION: NEW POLYNUCLEOTIDES AND POLYPEPTIDES OF THE IFNalpha-2 GENE
; FILE REFERENCE: 021349/0010
; CURRENT APPLICATION NUMBER: US/10/087,325
; CURRENT FILING DATE: 2002-03-01
; PRIOR APPLICATION NUMBER: FR 0102843
; PRIOR FILING DATE: 2001-03-01
; NUMBER OF SEQ ID NOS: 26
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 15
; LENGTH: 20
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-087-325-15

Query Match      100.0%; Score 10; DB 13; Length 20;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
   |||||:|:|:|
Db 20 CTTCTCTTTT 11

RESULT 24
US-09-853-830-168/c
; Sequence 168, Application US/09853830
; Patent No. US20020107388A1
; GENERAL INFORMATION:
; APPLICANT: Vandenberg, Arthur A.
; TITLE OF INVENTION: Methods of Identifying and Monitoring
; TITLE OF INVENTION: Disease-Associated T Cells
; FILE REFERENCE: P-IM 4734
; CURRENT APPLICATION NUMBER: US/09/853,830
; CURRENT FILING DATE: 2001-09-18
; NUMBER OF SEQ ID NOS: 184
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 168
; LENGTH: 24
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-853-830-168
```

```
Query Match      100.0%; Score 10; DB 10; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
   |||||:|:|:|
Db 18 CTTCTCTTTT 9

RESULT 25
US-09-934-489A-46/c
; Sequence 46, Application US/09934489A
; Publication No. US20030108872A1
; GENERAL INFORMATION:
; APPLICANT: Sulavik, Mark
; APPLICANT: Ling, Losee Lucy
; APPLICANT: Opperman, Tlm
; APPLICANT: Molr, Don
; APPLICANT: Bunker, Christopher
; TITLE OF INVENTION: Genomics-Assisted Rapid Identification of Targets
; FILE REFERENCE: 032796-082
; CURRENT APPLICATION NUMBER: US/09/934,489A
; CURRENT FILING DATE: 2001-08-23
; PRIOR APPLICATION NUMBER: 2001-08-23,896
; PRIOR FILING DATE: 2000-08-23
; NUMBER OF SEQ ID NOS: 63
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 46
; LENGTH: 24
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: primer
US-09-934-489A-46

Query Match      100.0%; Score 10; DB 11; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
   |||||:|:|:|
Db 16 CTTCTCTTTT 7

RESULT 26
US-10-438-729-167/c
; Sequence 167, Application US/10438729
; Publication No. US20030190665A1
; GENERAL INFORMATION:
; APPLICANT: Vandenberg, Arthur
; TITLE OF INVENTION: METHODS OF SELECTING T CELL RECEPTOR V PEPTIDES FOR THERAPEUTIC
; FILE REFERENCE: 6915-65828
; CURRENT APPLICATION NUMBER: US/10/438,729
; CURRENT FILING DATE: 2003-05-14
; PRIOR APPLICATION NUMBER: 60/203,984
; PRIOR FILING DATE: 2000-05-12
; PRIOR APPLICATION NUMBER: 09/853,830
; PRIOR FILING DATE: 2001-05-10
; PRIOR APPLICATION NUMBER: 60/380,731
; PRIOR FILING DATE: 2002-05-14
; NUMBER OF SEQ ID NOS: 181
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 167
; LENGTH: 24
; TYPE: DNA
; ORGANISM: Artificial sequence
; FEATURE:
; OTHER INFORMATION: synthetic
US-10-438-729-167

Query Match      100.0%; Score 10; DB 12; Length 24;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
```

```
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 18 CTTCTCTTTT 9

RESULT 27
US-09-827-998-1102/c
; Sequence 1102, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1102
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1102

Query Match 100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16

RESULT 28
US-09-827-998-1103/c
; Sequence 1103, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1103
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1103

Query Match 100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 24 CTTCTCTTTT 15

RESULT 29
US-09-827-998-1104/c
```

```
; Sequence 1104, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1104
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1104

Query Match 100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 23 CTTCTCTTTT 14

RESULT 30
US-09-827-998-1105/c
; Sequence 1105, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 1105
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1105

Query Match 100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
Db 22 CTTCTCTTTT 13

RESULT 31
US-09-827-998-1106/c
; Sequence 1106, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
```



```
Db      17  CTTCTCTTTT 8

RESULT 36
US-09-827-998-1111/c
; Sequence 1111, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 1111
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1111

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  CTTCTCTTTT 10
        |||||
Db      16  CTTCTCTTTT 7

RESULT 37
US-09-827-998-1112/c
; Sequence 1112, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Shannon, Mark
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 1112
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1112

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  CTTCTCTTTT 10
        |||||
Db      15  CTTCTCTTTT 6

RESULT 38
US-09-827-998-1113/c
; Sequence 1113, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
```

```
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 1113
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1113

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  CTTCTCTTTT 10
        |||||
Db      14  CTTCTCTTTT 5

RESULT 39
US-09-827-998-1114/c
; Sequence 1114, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Acomica Sequence Listing Engine
; SEQ ID NO 1114
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1114

Query Match      100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1  CTTCTCTTTT 10
        |||||
Db      13  CTTCTCTTTT 4

RESULT 40
US-09-827-998-1115/c
; Sequence 1115, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; APPLICANT: Shannon, Mark
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; CURRENT FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
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; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aescmca Sequence Listing Engine
; SEQ ID NO 1115
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1115

Query Match 100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 12 CTTCTCTTTT 3

RESULT 41
US-09-827-998-1116/c
; Sequence 1116, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aescmca Sequence Listing Engine
; SEQ ID NO 1116
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1116

Query Match 100.0%; Score 10; DB 10; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 11 CTTCTCTTTT 2

RESULT 42
US-09-827-998-1117/c
; Sequence 1117, Application US/09827998
; Patent No. US20020102252A1
; GENERAL INFORMATION:
; APPLICANT: Gu, Yizhong
; TITLE OF INVENTION: NOVEL ISOFORMS OF HUMAN PREGNANCY-ASSOCIATED PROTEIN E
; FILE REFERENCE: MDMORF-8
; CURRENT APPLICATION NUMBER: US/09/827,998
; PRIOR FILING DATE: 2001-04-06
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 1881
; SOFTWARE: Aescmca Sequence Listing Engine
; SEQ ID NO 1117
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-827-998-1117

Query Match 100.0%; Score 10; DB 10; Length 25;

Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 10 CTTCTCTTTT 1

RESULT 43
US-10-060-756A-3717/c
; Sequence 3717, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aescmca Sequence Listing Engine
; SEQ ID NO 3717
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3717

Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 25 CTTCTCTTTT 16

RESULT 44
US-10-060-756A-3718/c
; Sequence 3718, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30

```
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 3718
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3718
```

```
Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
Qy      1 CTTCTCTTTT 10
        |||||
Db      24 CTTCTCTTTT 15
```

```
RESULT 45
US-10-060-756A-3719/c
; Sequence 3719, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 3719
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3719
```

```
Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
Qy      1 CTTCTCTTTT 10
        |||||
Db      23 CTTCTCTTTT 14
```

```
RESULT 46
US-10-060-756A-3720/c
; Sequence 3720, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
```

```
Query Match      100.0%; Score 10; DB 14; Length 25;
```

```
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 3720
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3720
```

```
Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
Qy      1 CTTCTCTTTT 10
        |||||
Db      22 CTTCTCTTTT 13
```

```
RESULT 47
US-10-060-756A-3721/c
; Sequence 3721, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine
; SEQ ID NO 3721
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3721
```

```
Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
Qy      1 CTTCTCTTTT 10
```

Db 21 CTTCTCTTTT 12
|||||

RESULT 48

US-10-060-756A-3722/c
; Sequence 3722, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:

; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177

; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23

; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09

; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine

; SEQ ID NO 3722

; LENGTH: 25
; TYPE: DNA

; ORGANISM: Homo sapiens

US-10-060-756A-3722

Query Match 100.0%; Score 10; DB 14; Length 25;

Best Local Similarity 100.0%; Pred. No. 1.6e+04;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||

Db 20 CTTCTCTTTT 11
|||||

RESULT 49

US-10-060-756A-3723/c

; Sequence 3723, Application US/10060756A
; Publication No. US20030046717A1

; GENERAL INFORMATION:

; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN

; FILE REFERENCE: PB0177

; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23

; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09

; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine

; SEQ ID NO 3723

; LENGTH: 25
; TYPE: DNA

; ORGANISM: Homo sapiens

US-10-060-756A-3723

Query Match

Best Local Similarity 100.0%; Score 10; DB 14; Length 25;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||

Db 19 CTTCTCTTTT 10
|||||

RESULT 50

US-10-060-756A-3724/c

; Sequence 3724, Application US/10060756A
; Publication No. US20030046717A1

; GENERAL INFORMATION:

; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN

; FILE REFERENCE: PB0177

; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23

; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09

; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecomica Sequence Listing Engine

; SEQ ID NO 3724

; LENGTH: 25
; TYPE: DNA

; ORGANISM: Homo sapiens

US-10-060-756A-3724

Query Match

Best Local Similarity 100.0%; Score 10; DB 14; Length 25;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|||||

Db 18 CTTCTCTTTT 9
|||||

RESULT 51

US-10-060-756A-3725/c

; Sequence 3725, Application US/10060756A
; Publication No. US20030046717A1

; GENERAL INFORMATION:

; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN

; FILE REFERENCE: PB0177

; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-10-09

```
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00669
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00665
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00668
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00663
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: US 09/864,761
/ PRIOR FILING DATE: 2001-05-23
/ PRIOR APPLICATION NUMBER: US 60/327,898
/ PRIOR FILING DATE: 2001-10-09
/ NUMBER OF SEQ ID NOS: 4804
/ SOFTWARE: Acomica Sequence Listing Engine
/ SEQ ID NO 3725
/ LENGTH: 25
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-10-060-756A-3725
```

```
Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY 1 CTTCTCTTTT 10
    |||||
Db 17 CTTCTCTTTT 8
```

```
RESULT 52
US-10-060-756A-3726/c
/ Sequence 3726, Application US/10060756A
/ Publication No. US20030046717A1
/ GENERAL INFORMATION:
/ APPLICANT: Zhang, Jian
/ TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
/ FILE REFERENCE: PB0177
/ CURRENT APPLICATION NUMBER: US/10/060,756A
/ CURRENT FILING DATE: 2002-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00667
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00664
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00669
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00665
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00668
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00663
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: US 09/864,761
/ PRIOR FILING DATE: 2001-05-23
/ PRIOR APPLICATION NUMBER: US 60/327,898
/ PRIOR FILING DATE: 2001-10-09
/ NUMBER OF SEQ ID NOS: 4804
/ SOFTWARE: Acomica Sequence Listing Engine
/ SEQ ID NO 3726
/ LENGTH: 25
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-10-060-756A-3726
```

```
Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY 1 CTTCTCTTTT 10
    |||||
Db 16 CTTCTCTTTT 7
```

```
RESULT 53
US-10-060-756A-3727/c
/ Sequence 3727, Application US/10060756A
/ Publication No. US20030046717A1
/ GENERAL INFORMATION:
/ APPLICANT: Zhang, Jian
/ TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
/ FILE REFERENCE: PB0177
/ CURRENT APPLICATION NUMBER: US/10/060,756A
/ CURRENT FILING DATE: 2002-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00667
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00664
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00669
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00665
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00668
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00663
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: US 09/864,761
/ PRIOR FILING DATE: 2001-05-23
/ PRIOR APPLICATION NUMBER: US 60/327,898
/ PRIOR FILING DATE: 2001-10-09
/ NUMBER OF SEQ ID NOS: 4804
/ SOFTWARE: Acomica Sequence Listing Engine
/ SEQ ID NO 3727
/ LENGTH: 25
/ TYPE: DNA
/ ORGANISM: Homo sapiens
US-10-060-756A-3727
```

```
Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY 1 CTTCTCTTTT 10
    |||||
Db 15 CTTCTCTTTT 6
```

```
RESULT 54
US-10-060-756A-3728/c
/ Sequence 3728, Application US/10060756A
/ Publication No. US20030046717A1
/ GENERAL INFORMATION:
/ APPLICANT: Zhang, Jian
/ TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
/ FILE REFERENCE: PB0177
/ CURRENT APPLICATION NUMBER: US/10/060,756A
/ CURRENT FILING DATE: 2002-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00667
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00664
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00669
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00665
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00668
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: PCT/US01/00663
/ PRIOR FILING DATE: 2001-01-30
/ PRIOR APPLICATION NUMBER: US 09/864,761
/ PRIOR FILING DATE: 2001-05-23
/ PRIOR APPLICATION NUMBER: US 60/327,898
/ PRIOR FILING DATE: 2001-10-09
/ NUMBER OF SEQ ID NOS: 4804
/ SOFTWARE: Acomica Sequence Listing Engine
/ SEQ ID NO 3728
/ LENGTH: 25
```

```
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3728

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 14 CTTCTCTTTT 5

RESULT 55
US-10-060-756A-3729/c
; Sequence 3729, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecmica Sequence Listing Engine
; SEQ ID NO 3729
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3729

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

RESULT 56
US-10-060-756A-3730/c
; Sequence 3730, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecmica Sequence Listing Engine
; SEQ ID NO 3729
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3729

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 13 CTTCTCTTTT 4

RESULT 57
US-10-060-756A-3731/c
; Sequence 3731, Application US/10060756A
; Publication No. US20030046717A1
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aecmica Sequence Listing Engine
; SEQ ID NO 3731
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3731

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 11 CTTCTCTTTT 2

RESULT 58
US-10-060-756A-3732/c
; Sequence 3732, Application US/10060756A
; Publication No. US20030046717A1
```

```
; GENERAL INFORMATION:
; APPLICANT: Zhang, Jian
; TITLE OF INVENTION: HUMAN TESTIS EXPRESSED PATCHED LIKE PROTEIN
; FILE REFERENCE: PB0177
; CURRENT APPLICATION NUMBER: US/10/060,756A
; CURRENT FILING DATE: 2002-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 09/864,761
; PRIOR FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/327,898
; PRIOR FILING DATE: 2001-10-09
; NUMBER OF SEQ ID NOS: 4804
; SOFTWARE: Aeomica Sequence Listing Engine
; SEQ ID NO 3732
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-060-756A-3732

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
DB      10 CTTCTCTTTT 1

RESULT 59
US-10-215-112-7249/c
; Sequence 7249, Application US/10215112
; Publication No. US20030082596A1
; GENERAL INFORMATION:
; APPLICANT: Michael Mittmann
; TITLE OF INVENTION: Method of Genetic Analysis of Probes:
; FILE REFERENCE: Test3
; CURRENT APPLICATION NUMBER: US/10/215,112
; CURRENT FILING DATE: 2002-08-08
; NUMBER OF SEQ ID NOS: 14936
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 7249
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Oligonucleotide
US-10-215-112-7249

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
DB      13 CTTCTCTTTT 4

RESULT 60
US-10-215-112-7250/c
; Sequence 7250, Application US/10215112
; Publication No. US20030082596A1
```

```
; GENERAL INFORMATION:
; APPLICANT: Michael Mittmann
; TITLE OF INVENTION: Method of Genetic Analysis of Probes:
; FILE REFERENCE: Test3
; CURRENT APPLICATION NUMBER: US/10/215,112
; CURRENT FILING DATE: 2002-08-08
; NUMBER OF SEQ ID NOS: 14936
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 7250
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Oligonucleotide
US-10-215-112-7250

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
DB      12 CTTCTCTTTT 3

RESULT 61
US-10-215-112-7372/c
; Sequence 7372, Application US/10215112
; Publication No. US20030082596A1
; GENERAL INFORMATION:
; APPLICANT: Michael Mittmann
; TITLE OF INVENTION: Method of Genetic Analysis of Probes:
; FILE REFERENCE: Test3
; CURRENT APPLICATION NUMBER: US/10/215,112
; CURRENT FILING DATE: 2002-08-08
; NUMBER OF SEQ ID NOS: 14936
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 7372
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Oligonucleotide
US-10-215-112-7372

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CTTCTCTTTT 10
DB      12 CTTCTCTTTT 3

RESULT 62
US-10-098-263B-6553
; Sequence 6553, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:
; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; CURRENT FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 6553
; LENGTH: 25
; TYPE: DNA
```

```
; ORGANISM: Homo sapien
US-10-098-263B-6553

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 7 CTTCTCTTTT 16

RESULT 63
US-10-098-263B-10333
; Sequence 10333, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:
; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; CURRENT FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 10333
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-098-263B-10333

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 5 CTTCTCTTTT 14

RESULT 64
US-10-098-263B-34049
; Sequence 34049, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:
; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; CURRENT FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 34049
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-098-263B-34049

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 11 CTTCTCTTTT 20

RESULT 65
US-10-098-263B-41737/c
; Sequence 41737, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:
; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; CURRENT FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 41737
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-098-263B-41737

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 14 CTTCTCTTTT 5

RESULT 67
US-10-098-263B-114427
; Sequence 114427, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:
; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; CURRENT FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 114427
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-098-263B-86720/c
; Sequence 86720, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:
; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; CURRENT FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 86720
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-098-263B-86720

Query Match      100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
Db 14 CTTCTCTTTT 5
```


US-10-098-263B-114427

Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 16 CTTCTCTTTT 25

RESULT 68

US-10-098-263B-114428
; Sequence 114428, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:

; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; PRIOR FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 114428
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien

US-10-098-263B-114428

Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 16 CTTCTCTTTT 25

RESULT 69

US-10-098-263B-118057
; Sequence 118057, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:

; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; PRIOR FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 118057
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien

US-10-098-263B-118057

Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 15 CTTCTCTTTT 24

RESULT 70

US-10-098-263B-118058
; Sequence 118058, Application US/10098263B
; Publication No. US20030104410A1

; GENERAL INFORMATION:

; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; PRIOR FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 118058
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-098-263B-118058

Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 15 CTTCTCTTTT 24

RESULT 71

US-10-098-263B-118243
; Sequence 118243, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:

; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; PRIOR FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 118243
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien

US-10-098-263B-118243

Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 15 CTTCTCTTTT 24

RESULT 72

US-10-098-263B-118244
; Sequence 118244, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:

; APPLICANT: Mittman, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; PRIOR FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 118244
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien

US-10-098-263B-118244

Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 15 CTTCTCTTTT 24

RESULT 73

US-10-098-263B-119613
; Sequence 119613, Application US/10098263B
; Publication No. US20030104410A1
; GENERAL INFORMATION:
; APPLICANT: Mitten, Michael
; TITLE OF INVENTION: Human Microarray
; FILE REFERENCE: 3118.1
; CURRENT APPLICATION NUMBER: US/10/098,263B
; CURRENT FILING DATE: 2003-01-08
; PRIOR APPLICATION NUMBER: 60/276,759
; PRIOR FILING DATE: 2001-03-16
; NUMBER OF SEQ ID NOS: 131066
; SOFTWARE: Microarray Probe Sequence Listing Generator V 1.1
; SEQ ID NO 119613
; LENGTH: 25
; TYPE: DNA
; ORGANISM: Homo sapien
US-10-098-263B-119613

Query Match 100.0%; Score 10; DB 14; Length 25;
Best Local Similarity 100.0%; Pred. No. 1.6e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 13 CTTCTCTTTT 22

RESULT 74

US-09-765-272-359/c
; Sequence 359, Application US/09765272
; Patent No. US20020061545A1
; GENERAL INFORMATION:
; APPLICANT: Choi et. al.
; TITLE OF INVENTION: Streptococcus pneumoniae Antigens and Vaccines
; NUMBER OF SEQUENCES: 452
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: Maryland
; COUNTRY: USA
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage
; COMPUTER: HP Vectra 486/33
; OPERATING SYSTEM: MSDOS version 6.2
; SOFTWARE: ASCII Text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/765,272
; FILING DATE: 22-Jan-2001
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/961,083
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Brookes, A. Anders
; REGISTRATION NUMBER: 36,373
; REFERENCE/DOCKET NUMBER: PB340P2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (301) 309-8504
; TELEFAX: (301) 309-8512

; INFORMATION FOR SEQ ID NO: 359:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 33 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 359:
US-09-765-272-359

Query Match 100.0%; Score 10; DB 9; Length 33;
Best Local Similarity 100.0%; Pred. No. 1.5e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 23 CTTCTCTTTT 14

RESULT 75

US-09-780-533A-2783/c
; Sequence 2783, Application US/09780533A
; Publication No. US20030060611A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: Blatt, Larry
; APPLICANT: McSwiggen, Jim
; APPLICANT: Chowrira, Bharat
; APPLICANT: Haerberli, Pete
; TITLE OF INVENTION: Method and Reagent for the Inhibition of NOGO Gene
; FILE REFERENCE: MBH00.878-A (400/011)
; CURRENT APPLICATION NUMBER: US/09/780,533A
; CURRENT FILING DATE: 2001-02-09
; PRIOR APPLICATION NUMBER: US 60/181,797
; PRIOR FILING DATE: 2000-02-11
; NUMBER OF SEQ ID NOS: 6679
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 2783
; LENGTH: 38
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
US-09-780-533A-2783

Query Match 100.0%; Score 10; DB 11; Length 38;
Best Local Similarity 100.0%; Pred. No. 1.5e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
| | | | |
Db 38 CTTCTCTTTT 29

RESULT 76

US-09-780-533A-4693
; Sequence 4693, Application US/09780533A
; Publication No. US20030060611A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: Blatt, Larry
; APPLICANT: McSwiggen, Jim
; APPLICANT: Chowrira, Bharat
; APPLICANT: Haerberli, Pete
; TITLE OF INVENTION: Method and Reagent for the Inhibition of NOGO Gene
; FILE REFERENCE: MBH00.878-A (400/011)
; CURRENT APPLICATION NUMBER: US/09/780,533A
; CURRENT FILING DATE: 2001-02-09
; PRIOR APPLICATION NUMBER: US 60/181,797
; PRIOR FILING DATE: 2000-02-11
; NUMBER OF SEQ ID NOS: 6679
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 4693
; LENGTH: 38

; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
US-09-780-533A-4693

Query Match 100.0%; Score 10; DB 11; Length 38;
Best Local Similarity 30.0%; Pred. No. 1.5e+04;
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|::|::|::|::|
Db 29 CUUCUCUUU 38

RESULT 77

US-09-877-478-4478
; Sequence 4478, Application US/09877478
; Publication No. US2003068301A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: Draper, Kenneth
; APPLICANT: Blatt, Larry
; APPLICANT: McSwiggen, Jim
; APPLICANT: Morrissey, Dave
; TITLE OF INVENTION: Method and Reagent for Inhibiting Hepatitis B Virus Replication
; FILE REFERENCE: MBH00-845-H (400/029)
; CURRENT APPLICATION NUMBER: US/09/877,478
; CURRENT FILING DATE: 2001-12-31
; PRIOR APPLICATION NUMBER: US 07/882,712
; PRIOR FILING DATE: 1992-05-14
; PRIOR APPLICATION NUMBER: US 09/531,025
; PRIOR FILING DATE: 2000-03-20
; PRIOR APPLICATION NUMBER: US 09/636,385
; PRIOR FILING DATE: 2000-08-09
; PRIOR APPLICATION NUMBER: US 09/696,347
; PRIOR FILING DATE: 2000-10-24
; PRIOR APPLICATION NUMBER: US 08/193,627
; PRIOR FILING DATE: 1994-02-07
; PRIOR APPLICATION NUMBER: US 08/433,993
; PRIOR FILING DATE: 1995-05-04
; PRIOR APPLICATION NUMBER: US 08/434,504
; PRIOR FILING DATE: 1995-05-04
; PRIOR APPLICATION NUMBER: US 09/436,430
; PRIOR FILING DATE: 1999-11-08
; NUMBER OF SEQ ID NOS: 6586
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 4478
; LENGTH: 38
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (11)..(16)
; OTHER INFORMATION: Phosphorothioate 3'-Internucleotide Linkage
; NAME/KEY: misc feature
; LOCATION: (20)..(20)
; OTHER INFORMATION: Phosphorothioate 3'-Internucleotide Linkage
; NAME/KEY: misc feature
; LOCATION: (22)..(22)
; OTHER INFORMATION: Phosphorothioate 3'-Internucleotide Linkage
; NAME/KEY: misc feature
; LOCATION: (24)..(25)
; OTHER INFORMATION: Phosphorothioate 3'-Internucleotide Linkage
; NAME/KEY: misc feature
; LOCATION: (28)..(28)
; OTHER INFORMATION: Phosphorothioate 3'-Internucleotide Linkage
; NAME/KEY: misc feature
; LOCATION: (30)..(30)
; OTHER INFORMATION: Phosphorothioate 3'-Internucleotide Linkage
US-09-877-478-4478

Query Match 100.0%; Score 10; DB 11; Length 38;
Best Local Similarity 30.0%; Pred. No. 1.5e+04;
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|::|::|::|::|
Db 29 CUUCUCUUU 38

RESULT 78

US-09-792-818-944/c
; Sequence 944, Application US/09792818
; Publication No. US20030134806A1
; GENERAL INFORMATION:
; APPLICANT: Ribozyme Pharmaceuticals, Inc.
; APPLICANT: Jarvis, Thale
; APPLICANT: Von Carlowitz, Ira
; APPLICANT: McSwiggen, Jim
; APPLICANT: Hamblin, Paul
; APPLICANT: Ellis, Jonathan
; TITLE OF INVENTION: Method and Reagent for the Inhibition of Grb-2-related with Inse
; FILE REFERENCE: MBH00-901-A (400/013)
; CURRENT APPLICATION NUMBER: US/09/792,818
; CURRENT FILING DATE: 2001-02-23
; NUMBER OF SEQ ID NOS: 2304
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 944
; LENGTH: 38
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Enzymatic Nucleic Acid
US-09-792-818-944

Query Match 100.0%; Score 10; DB 12; Length 38;
Best Local Similarity 100.0%; Pred. No. 1.5e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTCTTTT 10
|::|::|::|::|
Db 38 CTTCTCTTTT 29

RESULT 79

US-09-922-261-227/c
; Sequence 227, Application US/09922261
; Patent No. US20020111471A1
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Putnam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/922,261
; CURRENT FILING DATE: 2001-08-03
; PRIOR APPLICATION NUMBER: US/09/461,697
; PRIOR FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 227
; LENGTH: 42
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-922-261-227

Query Match 100.0%; Score 10; DB 10; Length 42;

```
Best Local Similarity 100.0%; Pred. No. 1.5e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 14 CTTCTCTTTT 5

RESULT 80
US-10-032-585-1923
; Sequence 1923, Application US/10032585
; Publication No. US20030180953A1
; GENERAL INFORMATION:
; APPLICANT: Terry, Roemer D.
; APPLICANT: Bo, Jiang
; APPLICANT: Charles, Boone
; APPLICANT: Howard, Bussey
; TITLE OF INVENTION: Gene Disruption Methodologies for Drug Target Discovery
; FILE REFERENCE: 10182-005-999
; CURRENT APPLICATION NUMBER: US/10/032,585
; CURRENT FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 8000
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1923
; LENGTH: 43
; TYPE: DNA
; ORGANISM: Candida albicans
US-10-032-585-1923

Query Match 100.0%; Score 10; DB 12; Length 43;
Best Local Similarity 100.0%; Pred. No. 1.5e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 12 CTTCTCTTTT 21

RESULT 81
US-09-922-261-225/c
; Sequence 225, Application US/09922261
; Patent No. US2002011471A1
; GENERAL INFORMATION:
; APPLICANT: COGENT NEUROSCIENCE, Inc.
; APPLICANT: Lo, Donald C.
; APPLICANT: Barney, Shawn
; APPLICANT: Thomas, Mary Beth
; APPLICANT: Portbury, Stuart D.
; APPLICANT: Puranam, Kasturi
; APPLICANT: Katz, Lawrence C.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR DIAGNOSING
; TITLE OF INVENTION: AND TREATING CONDITIONS, DISORDERS, OR DISEASES INVOLVING
; FILE REFERENCE: 10001-005-999
; CURRENT APPLICATION NUMBER: US/09/922,261
; CURRENT FILING DATE: 2001-08-03
; PRIOR APPLICATION NUMBER: US/09/461,697
; PRIOR FILING DATE: 1999-12-14
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 225
; LENGTH: 48
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-922-261-225

Query Match 100.0%; Score 10; DB 10; Length 48;
Best Local Similarity 100.0%; Pred. No. 1.5e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 20 CTTCTCTTTT 11
```

```
RESULT 82
US-09-997-931-40/c
; Sequence 40, Application US/09997931
; Publication No. US20030087241A1
; GENERAL INFORMATION:
; APPLICANT: University of Rochester
; APPLICANT: Kool, Eric
; TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND DNA
; FILE REFERENCE: 220.00010142
; CURRENT APPLICATION NUMBER: US/09/997,931
; CURRENT FILING DATE: 2001-11-30
; PRIOR APPLICATION NUMBER: US 09/569,344
; PRIOR FILING DATE: 2000-05-11
; PRIOR APPLICATION NUMBER: US 08/805,631
; PRIOR FILING DATE: 1997-02-26
; PRIOR APPLICATION NUMBER: US 08/393,439
; PRIOR FILING DATE: 1995-02-23
; PRIOR APPLICATION NUMBER: US 08/047,860
; PRIOR FILING DATE: 1993-04-15
; NUMBER OF SEQ ID NOS: 129
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 40
; LENGTH: 53
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: 53mer circle
US-09-997-931-40
```

```
Query Match 100.0%; Score 10; DB 11; Length 53;
Best Local Similarity 100.0%; Pred. No. 1.5e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 25 CTTCTCTTTT 16
```

```
RESULT 83
US-09-997-931-41
; Sequence 41, Application US/09997931
; Publication No. US20030087241A1
; GENERAL INFORMATION:
; APPLICANT: University of Rochester
; APPLICANT: Kool, Eric
; TITLE OF INVENTION: CIRCULAR DNA VECTORS FOR SYNTHESIS OF RNA AND DNA
; FILE REFERENCE: 220.00010142
; CURRENT APPLICATION NUMBER: US/09/997,931
; CURRENT FILING DATE: 2001-11-30
; PRIOR APPLICATION NUMBER: US 09/569,344
; PRIOR FILING DATE: 2000-05-11
; PRIOR APPLICATION NUMBER: US 08/805,631
; PRIOR FILING DATE: 1997-02-26
; PRIOR APPLICATION NUMBER: US 08/393,439
; PRIOR FILING DATE: 1995-02-23
; PRIOR APPLICATION NUMBER: US 08/047,860
; NUMBER OF SEQ ID NOS: 129
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 41
; LENGTH: 53
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: stem-loop multimer which binds HIV-1 gag RNA
US-09-997-931-41
```

```
Query Match 100.0%; Score 10; DB 11; Length 53;
Best Local Similarity 30.0%; Pred. No. 1.5e+04;
Matches 3; Conservative 7; Mismatches 0; Indels 0; Gaps 0;
```

QY 1 CTTCTCTTTT 10
|::|::|::|
Db 38 CUUCUCUUU 47

RESULT 84
US-10-085-906-282
; Sequence 282, Application US/10085906
; Publication No. US20030054371A1
; GENERAL INFORMATION:
; APPLICANT: Ying, Vincent
; APPLICANT: Wu, Paul
; APPLICANT: Gray, Gary S.
; TITLE OF INVENTION: POLYMORPHIC ELEMENTS IN THE
; TITLE OF INVENTION: COSTIMULATORY RECEPTOR LOCUS AND USES THEREOF
; FILE REFERENCE: GNN-5343CP2
; CURRENT APPLICATION NUMBER: US/10/085,906
; CURRENT FILING DATE: 2002-02-27
; PRIOR APPLICATION NUMBER: US 60/126,215
; PRIOR FILING DATE: 1999-03-25
; PRIOR APPLICATION NUMBER: US 09/534,061
; PRIOR FILING DATE: 2000-03-24
; PRIOR APPLICATION NUMBER: PCT/US00/07938
; PRIOR FILING DATE: 2000-03-24
; NUMBER OF SEQ ID NOS: 545
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 282
; LENGTH: 59
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-085-906-282

Query Match 100.0%; Score 10; DB 14; Length 59;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
|::|::|::|
Db 45 CTTCTCTTTT 54

RESULT 85
US-09-908-975-18382
; Sequence 18382, Application US/09908975
; Publication No. US20030165843A1
; GENERAL INFORMATION:
; APPLICANT: SHOSHAN, Avi
; APPLICANT: WASSERMAN, Alon
; APPLICANT: MINTZ, Eli
; APPLICANT: MINTZ, Liat
; APPLICANT: FAIGLER, Simchon
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SPLICE
; TITLE OF INVENTION: THAT POPULATE A TRANSCRIPTOME
; FILE REFERENCE: 36688-0005
; CURRENT APPLICATION NUMBER: US/09/908,975
; CURRENT FILING DATE: 2001-07-20
; PRIOR APPLICATION NUMBER: US 60/287,724
; PRIOR FILING DATE: 2001-05-02
; PRIOR APPLICATION NUMBER: US 60/221,607
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 32337
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 18382
; LENGTH: 60
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-908-975-18382

Query Match 100.0%; Score 10; DB 12; Length 60;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10

Db 43 CTTCTCTTTT 52
|::|::|::|

RESULT 86
US-09-908-975-31961/c
; Sequence 31961, Application US/09908975
; Publication No. US20030165843A1
; GENERAL INFORMATION:
; APPLICANT: SHOSHAN, Avi
; APPLICANT: WASSERMAN, Alon
; APPLICANT: MINTZ, Eli
; APPLICANT: MINTZ, Liat
; APPLICANT: FAIGLER, Simchon
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SPLICE
; TITLE OF INVENTION: THAT POPULATE A TRANSCRIPTOME
; FILE REFERENCE: 36688-0005
; CURRENT APPLICATION NUMBER: US/09/908,975
; CURRENT FILING DATE: 2001-07-20
; PRIOR APPLICATION NUMBER: US 60/287,724
; PRIOR FILING DATE: 2001-05-02
; PRIOR APPLICATION NUMBER: US 60/221,607
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 32337
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 31961
; LENGTH: 60
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-908-975-31961

Query Match 100.0%; Score 10; DB 12; Length 60;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
|::|::|::|
Db 48 CTTCTCTTTT 39

RESULT 87
US-09-795-668-1332
; Sequence 1332, Application US/09795668
; Patent No. US20020045577A1
; GENERAL INFORMATION:
; APPLICANT: Stefansson, Hreinn
; APPLICANT: Steinhorsdottir, Valgerdur
; APPLICANT: Gulcher, Jeffrey R.
; TITLE OF INVENTION: HUMAN SCHIZOPHRENIA GENE
; FILE REFERENCE: 2345-2004-001
; CURRENT APPLICATION NUMBER: US/09/795,668
; CURRENT FILING DATE: 2001-02-28
; PRIOR APPLICATION NUMBER: US 09/515,716
; PRIOR FILING DATE: 2000-02-28
; NUMBER OF SEQ ID NOS: 1531
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1332
; LENGTH: 61
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-795-668-1332

Query Match 100.0%; Score 10; DB 9; Length 61;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTCTCTTTT 10
|::|::|::|
Db 17 CTTCTCTTTT 26

RESULT 88
US-09-795-686-1332

; Sequence 1332, Application US/09795686
; Patent No. US2002009495A1
; GENERAL INFORMATION:
; APPLICANT: Stefansson, Hreinn
; APPLICANT: Steinhorsdottir, Valgerdur
; APPLICANT: Gulcher, Jeffrey R.
; TITLE OF INVENTION: HUMAN SCHIZOPHRENIA GENE
; FILE REFERENCE: 2345.2005-001
; CURRENT APPLICATION NUMBER: US/09/795,686
; CURRENT FILING DATE: 2001-02-28
; PRIOR APPLICATION NUMBER: US/09/515,715
; PRIOR FILING DATE: 2000-02-28
; NUMBER OF SEQ ID NOS: 1531
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1332
; LENGTH: 61
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-795-686-1332

Query Match 100.0%; Score 10; DB 9; Length 61;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||
Db 17 CTTCTCTTTT 26

RESULT 89

US-09-946-807-1332
; Sequence 1332, Application US/09946807
; Patent No. US20020165144A1
; GENERAL INFORMATION:
; APPLICANT: Stefansson, Hreinn
; APPLICANT: Steinhorsdottir, Valgerdur
; APPLICANT: Gulcher, Jeffrey R.
; TITLE OF INVENTION: HUMAN SCHIZOPHRENIA GENE
; FILE REFERENCE: 2345.2004-001
; CURRENT APPLICATION NUMBER: US/09/946,807
; CURRENT FILING DATE: 2001-09-05
; PRIOR APPLICATION NUMBER: US/09/795,668
; PRIOR FILING DATE: 2001-02-28
; PRIOR APPLICATION NUMBER: US/09/515,716
; PRIOR FILING DATE: 2000-02-28
; NUMBER OF SEQ ID NOS: 1531
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1332
; LENGTH: 61
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-946-807-1332

Query Match 100.0%; Score 10; DB 10; Length 61;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||
Db 17 CTTCTCTTTT 26

RESULT 90

US-10-027-632-58348
; Sequence 58348, Application US/10027632
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; POLYMORPHISMS IN THE HUMAN GENOME
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006

; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 58348
; LENGTH: 61
; TYPE: DNA
; ORGANISM: Human
US-10-027-632-58348

Query Match 100.0%; Score 10; DB 13; Length 61;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||
Db 16 CTTCTCTTTT 25

RESULT 91

US-09-983-965-5096
; Sequence 5096, Application US/09983965
; Patent No. US20020137160A1
; GENERAL INFORMATION:
; APPLICANT: Warren, Wesley C.
; APPLICANT: Tao, Nengbing
; APPLICANT: Byatt, John C.
; APPLICANT: Mathialagan, Nagappan
; TITLE OF INVENTION: NUCLEIC ACID AND OTHER MOLECULES ASSOCIATED WITH LACTATION AND
; FILE REFERENCE: 37-21(10297)C
; CURRENT APPLICATION NUMBER: US/09/983,965
; CURRENT FILING DATE: 2001-10-26
; PRIOR APPLICATION NUMBER: US 09/465,231
; PRIOR FILING DATE: 1999-12-15
; PRIOR APPLICATION NUMBER: US 60/113,678
; PRIOR FILING DATE: 1998-12-17
; NUMBER OF SEQ ID NOS: 5912
; SEQ ID NO 5096
; LENGTH: 63
; TYPE: DNA
; ORGANISM: Bos taurus
; FEATURE:
; OTHER INFORMATION: Clone ID: 31-LIB34-084-Q1-E1-H11
US-09-983-965-5096

Query Match 100.0%; Score 10; DB 10; Length 63;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||
Db 24 CTTCTCTTTT 33

RESULT 92

US-10-085-906-276/c
; Sequence 276, Application US/10085906
; Publication No. US20030054371A1
; GENERAL INFORMATION:
; APPLICANT: Ying, Vincent
; APPLICANT: Wu, Paul

; APPLICANT: Gray, Gary S.
; TITLE OF INVENTION: POLYMORPHIC ELEMENTS IN THE
; FILE REFERENCE: GNN-5343CP2
; CURRENT APPLICATION NUMBER: US/10/085,906
; CURRENT FILING DATE: 2002-02-27
; PRIOR APPLICATION NUMBER: US 60/126,215
; PRIOR FILING DATE: 1999-03-25
; PRIOR APPLICATION NUMBER: US 09/534,061
; PRIOR FILING DATE: 2000-03-24
; PRIOR APPLICATION NUMBER: PCT/US00/07938
; PRIOR FILING DATE: 2000-03-24
; NUMBER OF SEQ ID NOS: 545
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 276
; LENGTH: 64
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-085-906-276

Query Match 100.0%; Score 10; DB 14; Length 64;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 43 CTTCTCTTTT 34

RESULT 93
US-09-908-975-27615
; Sequence 27615, Application US/09908975
; Publication No. US20030165843A1
; GENERAL INFORMATION:
; APPLICANT: SHOSHAN, Avi
; APPLICANT: WASSERMAN, Alon
; APPLICANT: MINTZ, Eli
; APPLICANT: MINTZ, Liat
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SPLICING
; FILE REFERENCE: 36688-0005
; CURRENT APPLICATION NUMBER: US/09/908,975
; CURRENT FILING DATE: 2001-07-20
; PRIOR APPLICATION NUMBER: US 60/287,724
; PRIOR FILING DATE: 2001-05-02
; PRIOR APPLICATION NUMBER: US 60/221,607
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 32337
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 27615
; LENGTH: 65
; TYPE: DNA
; ORGANISM: Mus musculus
US-09-908-975-27615

Query Match 100.0%; Score 10; DB 12; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 47 CTTCTCTTTT 56

RESULT 94
US-09-908-975-28061/c
; Sequence 28061, Application US/09908975
; Publication No. US20030165843A1
; GENERAL INFORMATION:
; APPLICANT: SHOSHAN, Avi
; APPLICANT: WASSERMAN, Alon
; APPLICANT: MINTZ, Eli

; APPLICANT: MINTZ, Liat
; APPLICANT: FAIGLER, Simchon
; TITLE OF INVENTION: OLIGONUCLEOTIDE LIBRARY FOR DETECTING RNA TRANSCRIPTS AND SPLICING
; FILE REFERENCE: 36688-0005
; CURRENT APPLICATION NUMBER: US/09/908,975
; CURRENT FILING DATE: 2001-07-20
; PRIOR APPLICATION NUMBER: US 60/287,724
; PRIOR FILING DATE: 2001-05-02
; PRIOR APPLICATION NUMBER: US 60/221,607
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 32337
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 28061
; LENGTH: 65
; TYPE: DNA
; ORGANISM: Mus musculus
US-09-908-975-28061

Query Match 100.0%; Score 10; DB 12; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 41 CTTCTCTTTT 32

RESULT 95
US-10-032-585-135
; Sequence 135, Application US/10032585
; Publication No. US20030180953A1
; GENERAL INFORMATION:
; APPLICANT: Terry, Roemer D.
; APPLICANT: Bo, Jiang
; APPLICANT: Charles, Boone
; APPLICANT: Howard, Bussey
; TITLE OF INVENTION: Gene Disruption Methodologies for Drug Target Discovery
; FILE REFERENCE: 10182-005-999
; CURRENT APPLICATION NUMBER: US/10/032,585
; CURRENT FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 8000
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 135
; LENGTH: 65
; TYPE: DNA
; ORGANISM: Candida albicans
US-10-032-585-135

Query Match 100.0%; Score 10; DB 12; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
Db 2 CTTCTCTTTT 11

RESULT 96
US-10-032-585-152
; Sequence 152, Application US/10032585
; Publication No. US20030180953A1
; GENERAL INFORMATION:
; APPLICANT: Terry, Roemer D.
; APPLICANT: Bo, Jiang
; APPLICANT: Charles, Boone
; APPLICANT: Howard, Bussey
; TITLE OF INVENTION: Gene Disruption Methodologies for Drug Target Discovery
; FILE REFERENCE: 10182-005-999
; CURRENT APPLICATION NUMBER: US/10/032,585
; CURRENT FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 8000
; SOFTWARE: PatentIn version 3.1

; SEQ ID NO 152
; LENGTH: 65
; TYPE: DNA
; ORGANISM: Candida albicans
US-10-032-585-152

Query Match 100.0%; Score 10; DB 12; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 15 CTTCTCTTTT 24

RESULT 97

US-10-032-585-2790
; Sequence 2790, Application US/10032585
; Publication No. US2003018953A1
; GENERAL INFORMATION:

; APPLICANT: Terry, Roemer D.
; APPLICANT: Bo, Jiang
; APPLICANT: Charles, Boone
; APPLICANT: Howard, Bussey
; TITLE OF INVENTION: Gene Disruption Methodologies for Drug Target Discovery
; FILE REFERENCE: 10182-005-999
; CURRENT APPLICATION NUMBER: US/10/032,585
; CURRENT FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 8000
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2790
; LENGTH: 65
; TYPE: DNA
; ORGANISM: Candida albicans
US-10-032-585-2790

Query Match 100.0%; Score 10; DB 12; Length 65;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 25 CTTCTCTTTT 34

RESULT 98

US-10-015-637-3
; Sequence 3, Application US/10015637
; Publication No. US20030046727A1
; GENERAL INFORMATION:

; APPLICANT: Wang, Qi
; APPLICANT: Dubois, Patrice
; APPLICANT: Liang, Jihong
; APPLICANT: Oulmassov, Tim
; TITLE OF INVENTION: Arcelin-5 Promoter and Uses Thereof
; FILE REFERENCE: 13587.106
; CURRENT APPLICATION NUMBER: US/10/015,637
; CURRENT FILING DATE: 2001-12-17
; PRIOR APPLICATION NUMBER: US 60/255879
; NUMBER OF SEQ ID NOS: 14
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 3
; LENGTH: 76
; TYPE: DNA
; ORGANISM: Glycine max
US-10-015-637-3

Query Match 100.0%; Score 10; DB 14; Length 76;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||

Db 43 CTTCTCTTTT 52
|||||

RESULT 99

US-10-029-386-24504
; Sequence 24504, Application US/10029386
; Publication No. US20030194704A1
; GENERAL INFORMATION:

; APPLICANT: Penn, Sharron G.
; APPLICANT: Hanzel, David K.
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; FILE REFERENCE: AEOMICA-X-2
; CURRENT APPLICATION NUMBER: US/10/029,386
; CURRENT FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 34288
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 24504
; LENGTH: 81
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO CHR18.3
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 1.1
; OTHER INFORMATION: EST_HUMAN HIT: AW905636.1, EVALUE 9.00e-02
US-10-029-386-24504

Query Match 100.0%; Score 10; DB 12; Length 81;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||
Db 29 CTTCTCTTTT 38

RESULT 100

US-10-029-386-17114/c
; Sequence 17114, Application US/10029386
; Publication No. US20030194704A1
; GENERAL INFORMATION:

; APPLICANT: Penn, Sharron G.
; APPLICANT: Hanzel, David K.
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; FILE REFERENCE: AEOMICA-X-2
; CURRENT APPLICATION NUMBER: US/10/029,386
; CURRENT FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 34288
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 17114
; LENGTH: 84
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AL137017.4
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.7
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 1.6
; OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 2.6
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.5
; OTHER INFORMATION: EST_HUMAN HIT: BE536158.1, EVALUE 9.50e-02
; OTHER INFORMATION: NT HIT: X03248.1, EVALUE 1.00e+00
US-10-029-386-17114

Query Match 100.0%; Score 10; DB 12; Length 84;
Best Local Similarity 100.0%; Pred. No. 1.4e+04;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CTTCTCTTTT 10
|||||

Wed Oct 29 15:38:01 2003

us-09-335-032-71.oli.rnpb

Page 29

Db 18 CTTCTTTT 9

Search completed: October 28, 2003, 18:49:54
Job time : 1754 secs